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(71) Applicant (for all designated States except US): INEX PHARMACEUTICALS CORPORATION [CA/CA]; 100-8900 Glenlyon Parkway, Glenlyon Business Park, Burnaby, British Columbia V5J 5J8 (CA).

(72) Inventors; and

(75) Inventors/Applicants (for US only): TAM, Ying, K.

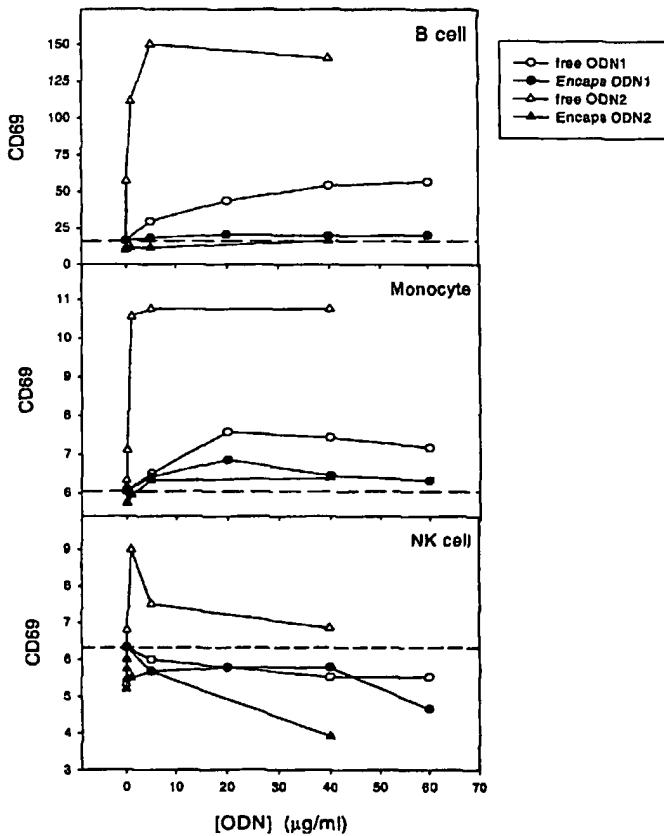
[CA/CA]; 3575 West 19th Avenue, Vancouver, British Columbia V6S 1C3 (CA). SEMPLE, Sean [CA/CA]; 2201-588 Broughton Street, Vancouver, British Columbia V6G 3E3 (CA). KLIMUK, Sandra [CA/CA]; 208-2190 West 7th Avenue, Vancouver, British Columbia V6K 4K7 (CA). CHIKH, Ghania [DZ/CA]; 1873 Nelson Street, Apt. #605, Vancouver, British Columbia, V6G 1M9 (CA).

(74) Agent: LONG, Erin, J.; Inex Pharmaceuticals Corporation, 100-8900 Glenlyon Parkway, Glenlyon Business Park, Burnaby, British Columbia V5J 5J8 (CA).

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[Continued on next page]

(54) Title: METHYLATED IMMUNOSTIMULATORY OLIGONUCLEOTIDES AND METHODS OF USING THE SAME



(57) Abstract: The invention discloses that methylated nucleic acids, particularly methylated oligonucleotides, and more particularly methylated oligonucleotides bearing a methylated cytosine of a CpG dinucleotide motif can be made immunostimulatory *in vivo*, by encapsulation of the nucleic acid in a lipid particle. It is further disclosed that encapsulated methylated nucleic acids that are ordinarily not immunostimulatory *in vivo* are as effective or even more effective than their encapsulated unmethylated counterparts. Also disclosed are methods for activating and/or expanding dendritic cell populations in response to antigenic stimulation using the compositions and methods disclosed herein.

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**METHYLATED IMMUNOSTIMULATORY OLIGONUCLEOTIDES
AND METHODS OF USING THE SAME**

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CROSS-REFERENCE TO RELATED APPLICATIONS

[001] This application claims priority to U.S. Provisional Patent Application Serial No. 60/379,343, filed May 10, 2002; and also to U.S. Patent Application Serial No. 10/290,545, filed 10 November 7, 2002; and also to U.S. Patent Application Serial No. 60/460,646 filed April 4, 2003.

TECHNICAL FIELD

15 [002] The invention relates to lipid-nucleic acid formulations and their methods of use in stimulating an immune response *in vivo*, and in particular, to liposomal formulations of nucleic acid sequences comprising at least one methylated cytosine so as to synergistically enhance their immunostimulatory activity, wherein the methylated cytosine preferably forms part of a CpG motif.

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BACKGROUND OF THE INVENTION

[003] Methylation of cytosine is the only known endogenous modification of DNA in eukaryotes, and occurs by the enzymatic addition of a methyl or hydroxymethyl group to the carbon-4 or 25 carbon-5 position of cytosine. Costello and Plass, 2001, *J. Med. Genet.* 38:285. The lower frequency of methylated cytosine residues found in bacterial DNA was suggested by many investigators as the reason why bacterial DNA could elicit an immune response *in vitro* and *in vivo*, in contrast to vertebrate DNA having a higher frequency of methylated cytosine residues which failed to stimulate any response. Messina *et al*, 1991, *J. Immunol* 147:1759. Later 30 studies using a peripheral blood mononuclear cell assay (PBMC) to measure mitogenicity of oligonucleotides *in vitro* showed that unmethylated DNA could be mitogenic but not methylated DNA. Of the many oligonucleotide sequences tested, those bearing a CpG dinucleotide motif were shown to be particularly mitogenic. Krieg *et al*, 1995, *Nature* 374:546-9. Oligonucleotides bearing a CpG dinucleotide motif were also shown to be immunostimulatory *in vivo*, provided 35 again, that the cytosine of the dinucleotide was not methylated. Parronchi P. *et al* 1999, *J. Immunol* 163:5946-53; Krieg A.M, 1999, *Biochim Biophys Acta* 1489:107-16.

[004] Similarly, U.S. Patent No 6,194,388 disclosed that B cell mitogenicity in T-cell depleted spleen cells was abolished when cytosines of the CpG motif were methylated but not when other cytosines were methylated, measuring mitogenicity based on *in vitro* thymidine incorporation into PBMC. Subsequently, based on screening over 300 oligonucleotide

5 sequences for their ability to induce B cell activation in T-cell depleted spleen cells measured by uridine uptake *in vitro*, related U.S. Patent No. 6,207,646 disclosed that oligonucleotides having unmethylated CpG dinucleotide motifs were more effective in stimulating mitogenicity than oligonucleotides lacking the CpG motif and that methylated counterparts of the same CpG oligonucleotides were not as effective. The site of methylation that most negatively impacted B-
10 cell mitogenicity was again shown to be the cytosine of the CpG dinucleotide, which when methylated reduced *in vitro* stimulation in comparison to the methylation of other cytosines.

[005] Similar conclusions were reached by the same group of applicants in U.S. Patent No. 6,239,116 (reduced NK cell lytic activity in methylated sequences), U.S. Patent No. 6,406,705

15 (non-CpG oligonucleotides lack adjuvant affect when combined with HBsAg), and U.S. Patent No. 6,429,199 (methylation of CpG motif caused a loss of stimulatory effect in combinations with GM-CSF). In general, therefore, methylated CpG oligonucleotides have been consistently shown to be either non-effective or much less effective than methylated oligonucleotides in stimulating mitogenic effects *in vitro* or in eliciting immunostimulatory effects *in vivo*.

20 [006] Recently, Schetter *et al.* in International Publication No. WO 02/069369 suggested that certain types of methylated CpG oligonucleotides may possess immunostimulatory activity based on an *in vitro* PBMC assay measuring production of certain leukocyte surface markers, including CD86, CD80, CD25 and CD69. The nucleic acid sequences tested, however, were heavily methylated at 4-9 sites in oligonucleotides having 1-4 CpG dinucleotides, and in all exemplified cases the methylated oligonucleotides remained less effective than their unmethylated counterparts. A variety of additional oligonucleotide structures were also investigated, including inosine substituted for guanosine, a cytosine adjacent to an inosine, a "dSpacer" having a sugar devoid of base substituted for a base adjacent to an inosine, a ZpG
25 dinucleotide where Z is replaced by 2-deoxyuridine, 5-fluoro-2'deoxy uridine, and a CpY dinucleotide where Y is a 2-aminopurine, xanthosine, N7-methyl-xanthosine, nebularine or a spacer. Unfortunately, no useful control sequence, such as a randomized sequence or mixture of sequences of the same length, was provided to demonstrate the predictive value of the *in vitro* assays and/or the accuracy of the applicants' conclusions with respect to the
30 proposed immunostimulatory activity that might occur *in vivo*. Moreover, no immunostimulatory activity was seen with dendritic cells, a key subset of antigen-presenting cells.

[007] There remains a continuing need in the art for improved adjuvant compositions having enhanced immunostimulatory activity. The compositions must be capable of stimulating a

wider range of antigen-presenting cells, including in particular dendritic cells. Further, the compositions must be capable, both alone and in combination with tumor, pathogen or other antigens, to stimulate effective immune responses *in vivo*.

5 [008] There is also an interest in increasing the breadth of nucleic acid sequence motifs that can be used as immunostimulatory adjuvants. What is needed, in particular, are compositions and methods capable of eliciting consistent immunostimulatory activity from CpG dinucleotide sequences having methylated cytosines, as well as other oligonucleotide structures demonstrating significant variability in their immunostimulatory activity *in vitro*.

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SUMMARY OF THE INVENTION

[009] The present inventors have discovered that the incorporation of methylated nucleic acid sequences into the lipid-nucleic acid (LNA) formulations described herein solves the 15 aforementioned problems in the prior art and provides synergistic benefits. In particular, when used in accordance with the present invention the immunostimulatory activity of methylated nucleic acid sequences can be significantly enhanced and improved immune stimulation consistently achieved *in vivo*, in marked contrast to the widely variable results reported in the prior art. Surprisingly, the lipid-methylated nucleic acid formulations of the present invention 20 also demonstrate therapeutic efficacy that is as good as, and in many cases better than, similar lipid-nucleic acid formulations employing the corresponding unmethylated sequences. Further, immunostimulatory activity is obtained in a broader class of antigen-presenting cells, and in particular in dendritic cells. Moreover, as demonstrated herein, and unlike the prior art, effective immune stimulation can be achieved with encapsulated nucleic acids having only a 25 single methylated CpG dinucleotide.

[0010] In one aspect, the invention provides lipid-methylated nucleic acid formulations for stimulating an immune response in an animal, comprising a lipid component and a nucleic acid component comprising at least one methylated nucleic acid sequence. In certain preferred 30 embodiments, the methylated nucleic acid sequence comprises a methyl or hydroxymethyl group attached to the carbon-4 position (4-mC) or carbon-5 position (5-mC) of at least one cytosine, wherein the methylated cytosine residue will generally be part of a CpG dinucleotide motif located in said sequence. In alternative preferred embodiments, the methylated nucleic acid is fully encapsulated by the lipid component to form a liposomal particle, as further 35 described herein.

[0011] In certain embodiments, the methylated nucleic acid sequence lacks immunostimulatory activity *in vivo* when administered to an animal as a free nucleic acid. In other embodiments, the lipid-methylated nucleic acid formulation is either equivalent to or more immunostimulatory

in vivo than a corresponding lipid-nucleic acid formulation having the same sequence but lacking methylation of one or more cytosine residues.

[0012] In one embodiment, the nucleic acid sequence comprises at least one CpG dinucleotide 5 having a methylated cytosine. In a preferred embodiment, the nucleic acid sequence comprises a single CpG dinucleotide, wherein the cytosine in said CpG dinucleotide is methylated. In a specific embodiment, the nucleic acid sequence comprises the sequence 5' TAACGTTGAGGGGCAT 3' (ODN1m). In an alternative embodiment, the nucleic acid sequence comprises at least two CpG dinucleotides, wherein at least one cytosine in the CpG 10 dinucleotides is methylated. In a further embodiment, each cytosine in the CpG dinucleotides present in the sequence is methylated. In another specific embodiment, the nucleic acid sequence comprises the sequence 5' TTCCATGACGTTCCCTGACGTT 3' (ODN2m). In another embodiment, the nucleic acid sequence comprises a plurality of CpG dinucleotides, wherein at least one of said CpG dinucleotides comprises a methylated cytosine. Significantly however, 15 and unlike the prior art teachings, effective immune stimulation may be obtained as described herein utilizing nucleic acid sequences having only a single CpG dinucleotide with a methylated cytosine, or a plurality of CpG dinucleotides wherein only one or a couple of the cytosines of said CpG dinucleotides are methylated.

20 [0013] In preferred embodiments, the lipid-methylated nucleic acid formulation is capable of activating and/or expanding dendritic cells when administered to an animal *in vivo*. In one aspect, dendritic cells bearing at least one of a CD11c and DEC205 marker are expanded *in vivo* upon administration of the subject formulations, preferably in conjunction with antigenic stimulation. In another aspect, dendritic cells bearing the CD11c or DEC205 marker are 25 activated *in vivo* to express a CD86 marker after administration of the subject formulations, again preferably in conjunction with antigenic stimulation.

[0014] In various embodiments, the lipid-nucleic acid formulation further comprises a 30 pharmaceutically acceptable carrier, buffer or diluent.

35 [0015] In certain embodiments, the nucleic acid is comprised of a phosphodiester backbone. In other embodiments, the nucleic acid is comprised of a non- phosphodiester backbone. In more particular embodiments, the non-phosphodiester backbone is a phosphorothioate backbone.

[0016] In various embodiments of the composition the liposomal particle comprises a cationic lipid. Example cationic lipids are selected from a group of cationic lipids consisting of DDAB, DODAC, DOTAP, DMRIE, DOSPA, DMDMA, DC-Chol, DODMA, DODAP and mixtures thereof. In further embodiments, the liposomal particle further comprises a neutral lipid selected from

the group consisting of DOPE, DSPC, POPC, diacylphosphatidylcholine, diacylphosphatidylethanolamine, ceramide, sphingomyelin, cephalin, cerebrosides, and mixtures thereof. In other embodiments, the liposomal particle alone comprises a neutral lipid selected from the group consisting of DOPE, DSPC, POPC, diacylphosphatidylcholine, 5 diacylphosphatidylethanolamine, ceramide, sphingomyelin, cephalin, cerebrosides and mixtures thereof. In other embodiments, the lipid particle comprises a lipid selected from the group consisting of but not limited to DODAP, DODMA, DSPC, POPC, and mixtures of thereof. In certain other embodiments, the lipid particle is comprised of a mixture of sphingomyelin and a lipid selected from the group consisting of DODAP, DODMA, DSPC, POPC, and mixtures of 10 thereof. In still other embodiments, the lipid component comprises DSPC, DODMA, Chol, and PEG-DMG and the ratio of said DSPC to said DODMA to said Chol to said PEG-DMG is about 20:25:45:10 mol/mol. In a further more specific embodiment, the ratio of said lipid component to said nucleic component is about 0.01-0.25 wt/wt.

15 [0017] In other embodiments the lipid particle further includes a steric barrier lipid component on the surface of the lipid particle. In certain embodiments, the steric barrier lipid component is selected from the group consisting of PEG-DMG, PEG-PE, and a PEG ceramide. In one embodiment, the PEG ceramide is PEG-ceramide C-14. In another embodiment the PEG ceramide is PEG-ceramide C-20.

20 [0018] In a still further embodiment, a composition is provided comprising an antigen of interest in combination with the aforementioned lipid-methylated nucleic acid formulations. The antigen may be either mixed with or associated with the lipid-methylated nucleic acid formulation. Preferably, the antigen is associated with the formulation, as described herein. In one 25 embodiment, the antigen is a tumor antigen. In preferred embodiments, the methylated nucleic acid sequence comprises a 4-mC or 5-mC located within at least one CpG dinucleotide motif. In particularly preferred embodiments, the methylated nucleic acid sequence is encapsulated in a liposomal particle. In further embodiments, the antigen is also encapsulated in the liposomal particle.

30 [0019] In another aspect, the invention provides methods of stimulating enhanced immune activity in an animal comprising administering any of the foregoing compositions to the animal in order to induce an improved immune response. The LNA formulations may be used directly as adjuvants, or may advantageously be combined with one or more target antigens in vaccine 35 formulations. Preferably, administration of the subject compositions is capable of stimulating one or more dendritic cells present in the animal's immune system. In one embodiment, the target antigen is administered in association with the lipid-nucleic acid formulations described herein, and more preferably with a liposomal particle. In a further preferred embodiment, the

antigen is encapsulated in the liposomal particle. In certain embodiments the antigen comprises one or more epitopes from one or more tumor antigens or microbial antigens.

[0020] In a further aspect, methods for stimulating antigen-presenting cells are provided,
5 comprising the step of contacting at least one antigen-presenting cell *in vitro, ex vivo or in vivo* with an immunostimulatory composition as described herein. In preferred embodiments, the antigen-presenting cell comprises a dendritic cell.

[0021] In one embodiment, methods for expanding dendritic cells *in vivo* are provided,
10 comprising administering to a host the subject lipid-nucleic acid formulations comprising a nucleic acid sequence having at least one 4mC or 5mC located within a CpG dinucleotide, wherein said administration is effective to expand dendritic cells in said host. Preferably, expansion of said dendritic cells is characterized by an increase in the number of host antigen-presenting cells expressing at least one of a CD11c and DEC205 marker.

15 [0022] In another embodiment, methods for activating dendritic cells *in vivo* are provided, comprising administering to a host the subject lipid-nucleic acid formulations comprising a nucleic acid sequence having at least one 4mC or 5mC located within a CpG dinucleotide, wherein said administration is effective to expand dendritic cells in said host in response to
20 antigenic stimulation. Preferably, activation of said dendritic cells is characterized by an increase in the number of host antigen-presenting cells co-expressing at least one of a CD11c or DEC205 marker in conjunction with a CD86 marker. In a particularly preferred embodiment, antigenic stimulation is achieved by administration of the subject formulations in combination with one or more target antigens of interest, either mixed with or associated with the lipid-methylated nucleic acid formulation.

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BRIEF DESCRIPTION OF THE DRAWINGS

[0023] Figure 1 illustrates *in vitro* stimulation of leukocytes bearing the activation marker CD69
30 results from treating whole blood with free oligonucleotides. Mouse whole blood was treated *in vitro* with either the free oligonucleotide herein designated ODN1 or with the oligonucleotide designated ODN2.

[0024] Figure 2 illustrates *in vivo* treatment of mice by injection with encapsulated or free ODN1
35 and ODN2 oligonucleotides produces results that are contrary to those obtained *in vitro*.

[0025] Figure 3 shows that when encapsulated in a lipid vesicle the methylated ODN1m was more active than the unmethylated counterpart ODN1 in stimulating activation of dendritic cells

in vivo.

[0026] Figure 4A shows that both the methylated ODN1m and the unmethylated ODN1 stimulated the expansion of CD11c positive cells in spleen and whole blood.

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[0027] Figure 4B shows that both ODN1 and ODN1m stimulate the expansion of DEC205 positive cells in spleen, whole blood and lymph node.

10 [0028] Figure 5 shows that the methylated ODN1m was more active than the unmethylated counterpart ODN1, in stimulating CD86 expression when either ODN was lipid encapsulated.

[0029] Figure 6 shows that *in vivo* administration of free oligonucleotide had no affect on stimulation of IL-6, IL-12 IFN-gamma or MCP-1. In contrast, *in vivo* administration of lipid encapsulated oligonucleotides stimulated production of each of these cytokines.

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[0030] Figure 7A illustrates increased IL-12 induction by treatment of mice with either encapsulated PO or PS oligonucleotide ODN1 in comparison to free oligonucleotide ODN1 measured over an oligonucleotide dosage scale. Figure 7B shows that treatment with encapsulated PO oligonucleotides stimulates a strong early induction of IFN-gamma while 20 treatment with encapsulated PS oligonucleotides stimulates a smaller but still effective induction of IFN-gamma.

[0031] Figure 8 shows a comparison of IgM titres indicative of a Th-1 response upon administration of free PS or PO oligonucleotides.

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[0032] Figure 9 shows a comparison of IgG production indicative of a Th-2 response upon administration of free PS or PO oligonucleotide, including methylated oligonucleotides

30 [0033] Figure 10 shows that over a series of screenings of animals treated with methylated or unmethylated lipid encapsulated oligonucleotides, the methylated oligonucleotides are about the same or better than the unmethylated oligonucleotide in stimulating proliferation of dendritic cells, NK cells and CD8+ T-cells .

35 [0034] Figures 11A and B show that over a series of screenings of animals treated with methylated or unmethylated lipid encapsulated oligonucleotides, the methylated oligonucleotides are better than the unmethylated oligonucleotide in stimulating proliferation of cytotoxic T lymphocytes and Ag-specific lymphocytes.

[0035] Figure 11C illustrates data from a representative tetramer study that was included in the overall screenings described in Figures 11A and 11B.

[0036] Figure 12 illustrates that when administered to an animal as free oligonucleotides, 5 methylated versions have less therapeutic efficacy than methylated nucleotides in reducing tumor growth.

[0037] Figure 13 illustrates that encapsulation of oligonucleotides provides improved efficacy of methylated and unmethylated oligonucleotides over free ODN, particularly when the 10 oligonucleotides contain a natural phosphorothioate (PS) backbone.

[0038] Figure 14 shows that encapsulation of oligonucleotides provides improved efficacy of methylated and unmethylated oligonucleotides over free ODN, when the oligonucleotides contain a phosphodiester (PO) backbone. 15

[0039] Figure 15 shows that lipid encapsulated PS oligonucleotides ODN2 and ODN2m each exhibit therapeutic efficacy.

[0040] Figure 16 illustrates an adjuvant effect and therapeutic efficacy of administering the 20 methylated ODN1m to an animal inoculated with a B16 melanoma tumor. Encapsulation of the ODN1m oligonucleotide in a lipid particle increased its efficacy in reducing tumor volume.

[0041] Figure 17 shows that for a series of mice inoculated with the B16 melanoma and subsequently treated by administration of a 20mg/kg dose of oligonucleotide, the average tumor 25 size of tumours in mice treated with encapsulated free oligonucleotides ODN1 and ODN1m.

[0042] Figure 18 shows the reduction in tumor volume when mice were treated with encapsulated methylated ODN1m and the unmethylated counterpart ODN1.

30 [0043] Figure 19 shows survival rates of mice treated with the encapsulated methylated ODN1m in comparison to treatment with the unmethylated ODN1 in two different studies.

[0044] Figure 20 illustrates the efficacy in terms of tumor volume when methylated ODN1m and the unmethylated counterpart ODN1 are encapsulated in a lipid particle.

35 [0045] Figure 21 shows the survival rate of mice treated with encapsulated methylated ODN1m relative to treatment with the unmethylated encapsulated ODN1.

[0046] Figure 22 illustrates that encapsulated PS oligonucleotides ODN1 and ODN2 produced an IFN-gamma peak that is not produced by encapsulated PO oligonucleotides 6 days after treatment.

5 [0047] Figure 23 shows the effect on blood clearance in mice methylated or unmethylated oligonucleotides encapsulated in lipid particles having different PEG-ceramide steric coatings.

10 [0048] Figure 24 illustrates therapeutic efficacy of liposomal particles encapsulating methylated or methylated CpG oligonucleotide in treating a tumor by administering the composition to an animal having the tumor.

[0049] Figure 25 illustrates that lipid encapsulation of methylated PS-ODN5m provided a more effective therapeutic benefit than encapsulation of the equivalent unmethylated PS-ODN5 at reducing tumor growth over time.

15 [0050] Figure 26 shows the survival rate of the mice treated with free and encapsulated methylated ODN5m relative to treatment with the unmethylated encapsulated and free ODN5.

20 [0051] Figure 27 illustrates efficacy in terms of tumor volume when treated with free unmethylated and methylated PS and PO ODN 7 and free and encapsulated PO-ODN7m.

[0052] Figure 28 shows survival rates of mice treated with the free unmethylated and methylated PS and PO ODN7 in comparison to treatment with the encapsulated PO ODN7m.

25 [0053] Figure 29 shows the CTL response to a B16 cell target after immunization with a multiple epitope cancer vaccine using encapsulated ODN 1m.

[0054] Figure 30 shows the CTL response to a B16 cell target after immunization with a multiple epitope cancer vaccine using peptide-pulsed dendritic cells.

30 [0055] Figure 31 shows the CTL response to a B16 cell target after immunization with tumor cell lysate in combination with encapsulated ODN 1m or dendritic cells.

DETAILED DESCRIPTION OF THE INVENTION

35 [0056] The invention provides formulations and methods of use thereof, based on the discovery that methylated nucleic acids, particularly methylated oligonucleotides, and more particularly methylated oligonucleotides bearing a methylated cytosine of a CpG dinucleotide motif can enhance stimulation of immune responses either *in vitro*, *ex vivo* and *in vivo*, by encapsulation

of the nucleic acid in a lipid particle. It is further disclosed that lipid-encapsulated methylated nucleic acids, which are ordinarily not immunostimulatory in their free form *in vivo*, can in fact be just as effective and in some cases more effective at stimulating immune responses when encapsulated in the subject formulations in comparison with their unmethylated counterparts.

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[0057] The invention is exemplified by testing methylated and unmethylated counterparts of various oligonucleotides, configured with various backbones and encapsulated in various formulations of lipid particles. The lipid encapsulated methylated oligonucleotides are immunostimulatory with ordinary phosphodiester (PO) backbones as well as phosphorothioate (PS) backbones. It is further disclosed that, in some cases, the PO backbones may enhance a Th-1 mediated cellular immune response, while the PS backbones may stimulate a Th-1 mediated humoral immune response. In certain aspects of the invention, the lipid encapsulated methylated oligonucleotides are further combined with target antigens, particularly microbial antigens and/or tumor-associated antigens.

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[0058] In particular, when used in accordance with the present invention the immunostimulatory activity of methylated nucleic acid sequences is significantly enhanced and improved immune stimulation is consistently achieved *in vivo*, in marked contrast to the widely variable results reported in the prior art. Further, immunostimulatory activity is obtained in a broader class of antigen-presenting cells, and in particular in dendritic cells. The invention demonstrates for the first time, activation and expansion of dendritic cells by treatment with methylated oligonucleotides when they are lipid encapsulated. Accordingly methods of enhancing activation and/or expansion of dendritic cells is another aspect of the invention. Detailed methods of making, using and testing the various formulations of the invention are described hereafter and in the references cited herein, all of which are incorporated by reference.

Abbreviations and Definitions

[0059] The following abbreviations are used herein: RBC, red blood cells; DDAB, N,N-distearyl-N,N-dimethylammonium bromide; DODAC, N,N-dioleyl-N,N-dimethylammonium chloride; DOPE, 1,2-sn-dioleoylphosphatidylethanolamine; DOSPA, 2,3-dioleyloxy-N-(2-sperminecarboxamido)ethyl)-N,N-dimethyl-1-propanaminium trifluoroacetate; DOTAP, 1,2-dioleyloxy-3-(N,N,N-trimethylamino)propane chloride; DOTMA, 1,2-dioleyloxy-3-(N,N,N-trimethylamino)propanechloride; OSDAC, N-oleyl-N-stearyl-N,N-dimethylammonium chloride; RT, room temperature; HEPES, 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid; FBS, fetal bovine serum; DMEM, Dulbecco's modified Eagle's medium; PEG-Cer-C.sub.14, 1-O-(2'-(.omega.-methoxypolyethyleneglycol)succinoyl)-2-N-myristoyl-sphing osine; PEG-Cer-C.sub.20, 1-O-(2'-(.omega.-methoxypolyethyleneglycol)succinoyl)-2-N-arachidoyl-sphing osine; PBS, phosphate-buffered saline; THF, tetrahydrofuran; EGTA, ethylenebis(oxyethylenenitrilo)-tetraacetic acid; SF-DMEM, serum-free DMEM; NP40, nonylphenoxypolyethoxyethanol, 1,2

dioleoyl-3 dimethylaminopropane (DODAP), palmitoyl oleoyl phosphatidylcholine (POPC) and distearoylphosphatidylcholine (DSPC).

[0060] The technical and scientific terms used herein have the meanings commonly understood by one of ordinary skill in the art to which the present invention pertains, unless otherwise defined. Reference is made herein to various methodologies known to those of skill in the art. Publications and other materials setting forth such known methodologies to which reference is made are incorporated herein by reference in their entirety as though set forth in full. Standard reference works setting forth the general principles of recombinant DNA technology include Sambrook, J., *et al.*, *Molecular Cloning: A Laboratory Manual*, 2d Ed., Cold Spring Harbor Laboratory Press, Planview, N.Y. (1989); McPherson, M. J., Ed., *Directed Mutagenesis: A Practical Approach*, IRL Press, Oxford (1991); Jones, J., *Amino Acid and Peptide Synthesis*, Oxford Science Publications, Oxford (1992); Austen, B. M. and Westwood, O. M. R., *Protein Targeting and Secretion*, IRL Press, Oxford (1991). Any suitable materials and/or methods known to those of skill can be utilized in carrying out the present invention; however, preferred materials and/or methods are described. Materials, reagents and the like to which reference is made in the following description and examples are obtainable from commercial sources, unless otherwise noted. It is believed that one skilled in the art can, based on the description herein, utilize the present invention to its fullest extent. The entire contents of all of the references (including literature references, issued patents, published patent applications, and co-pending patent applications) cited throughout this application are hereby expressly incorporated by reference.

[0061] The immunostimulatory compositions used in the methods of the present invention will generally be referred to as lipid-therapeutic agent ("LTA") formulations comprising at least one lipid component and at least one therapeutic agent, and having greater immunostimulatory activity than the therapeutic agent alone, *in vivo*. "Therapeutic agent" or "therapeutic compound" or "drug" as used herein can be used interchangeably and refer to any synthetic, recombinant, or naturally occurring molecule that provides a beneficial effect in medical treatment of a subject. Examples of therapeutic agents include, but are not limited to nucleic acids, peptides, and chemicals.

[0062] In the preferred embodiments described herein, the therapeutic agent comprises at least one methylated nucleic acid sequence, more preferably at least one methylated oligonucleotide, and most preferably at least one methylated oligodeoxynucleotide ("ODN"). In preferred embodiments, the methylated cytosine residue is part of a CpG dinucleotide motif located in said sequence. The CpG comprises a methyl or hydroxymethyl group attached to the carbon-4 position (4-mC) or carbon-5 position (5-mC) of at least one cytosine. In further embodiments, the methylated nucleic acid sequence may alternatively or additionally comprise

methyl modifications of the deoxyribose or ribose sugar moiety as described in Henry *et al.* 2000 J. Pharmacol. Exp. Ther. 292:468, Zhao *et al.* 1999 Bioorg. Med. Chem Lett. 9:3453, Zhao *et al.* 2000 Bioorg Med. Chem Lett. 10:1051. In a particularly preferred embodiment, the ODN comprises a methylated nucleic acid sequence that has immunostimulatory activity and is 5 designated an immunostimulatory sequence ("ISS") in non-methylated form.

[0063] "Subject" or "host" as used herein refers to an organism, male or female, having an immune system, preferably an animal, more preferably a vertebrate, even more preferably a mammal, still even more preferably a rodent, and most preferably a human. Further examples 10 of a subject include, but are not limited to, dogs, cats, cows, horses, pigs, sheep, goats, mice, rabbits, and rats. "Patient" as used herein refers to a subject in need of treatment for a medical condition (e.g., disease or disorder).

[0064] "*In vivo*" as used herein refers to an organism, preferably in a mammal, more preferably 15 in a rodent, and most preferably in a human.

[0065] "Immunostimulatory," "immunostimulatory activity" or "stimulating an immune response," and grammatical equivalents thereof, as used herein refers to inducing, increasing, enhancing, or modulating an immune response, or otherwise providing a beneficial effect with respect to an 20 immune response. As used herein "immune response" refers to systemic and/or mucosal immune responses. Preferably, and in view of the wide variation in *in vitro* experimental results reported in the prior art, the immunostimulatory activity of a given formulation and nucleic acid sequence is determined in a suitable *in vivo* assay as described herein.

[0066] "A target antigen" as used herein refers to an antigen of interest to which a immune 25 response can be directed or stimulated. The target antigen used in the compositions of the present invention for stimulating an immune response directed to that target antigen may be a synthetic, naturally-occurring or isolated molecule or a fragment thereof, and may comprise single or multiple epitopes. Thus, the compositions of the present invention may stimulate 30 immune responses directed to single or multiple epitopes of an antigen. In preferred embodiments, the target antigen is associated with the lipid particles of the present invention. "In association with", "associated with", or grammatical equivalents thereof, as used herein with 35 reference to an antigen (or target antigens), refers to antigens that are attached to or encapsulated by another component. With reference to the lipid particles or liposomes of the present invention, the antigen may be, for example, encapsulated in the lumen or intralamellar spaces of the lipid particles; disposed or attached within or partially within the lipid membrane, or attached (e.g., covalently or ionically) to the lipid particle. The antigen may be attached to the interior of the lipid particle or, more preferably, the antigen is attached to the exterior of the

lipid particle. In preferred embodiments the antigen is encapsulated within the lipid particle.

[0067] Examples of antigens useful in the compositions and methods of the present invention include, but are not limited to, peptides or proteins, cells, cell extracts, polysaccharides, 5 polysaccharide conjugates, lipids, glycolipids, glycopeptides, and carbohydrates. In one embodiment, the antigen is in the form of a peptide or protein antigen. In another embodiment, the antigen is a nucleic acid encoding a peptide or protein in a form suitable for expression in a subject and presentation to the immune system of that subject. In a preferred embodiment, the compositions used in the methods of the present invention comprise a peptide or protein target 10 antigen that stimulates an immune response to that target antigen in a mammal. Preferably, the target antigen is a pathogen ("target pathogen") capable of infecting a mammal including, for example, bacteria, viruses, fungi, yeast, parasites and other microorganisms capable of infecting mammalian species. Alternatively, the target antigen may be a tumor-associated antigen.

15 [0068] A "tumor-associated antigen" as used herein is a molecule or compound (e.g., a protein, peptide, polypeptide, lipid, glycolipid, carbohydrate and/or DNA) associated with a tumor or cancer cell and which is capable of provoking an immune response when expressed on the surface of an antigen presenting cell in the context of an MHC molecule. Tumor-associated 20 antigens include self antigens, as well as other antigens that may not be specifically associated with a cancer but nonetheless enhance an immune response to and/or reduce the growth of a cancer when administered to an animal. In view of the potential risk of autoimmune reactions, the use of self antigens in the subject vaccines may be limited to non-critical tissues such as breast, prostate, testis, melanocytes, etc. More specific embodiments are provided herein.

25 [0069] A "microbial antigen" as used herein is an antigen of a microorganism and includes but is not limited to, infectious virus, infectious bacteria, infectious parasites and infectious fungi. Microbial antigens may be intact microorganisms, and natural isolates, fragments, or derivatives thereof, synthetic compounds which are identical to or similar to naturally-occurring 30 microbial antigens and, preferably, induce an immune response specific for the corresponding microorganism (from which the naturally-occurring microbial antigen originated). In a preferred embodiment, a compound is similar to a naturally-occurring microorganism antigen if it induces an immune response (humoral and/or cellular) to a naturally-occurring microorganism antigen. Compounds or antigens that are similar to a naturally-occurring microorganism antigen are well 35 known to those of ordinary skill in the art. A non-limiting example of a compound that is similar to a naturally-occurring microorganism antigen is a peptide mimic of a polysaccharide antigen. More specific embodiments are provided herein.

[0070] The term "antigen" is further intended to encompass peptide or protein analogs of known or wild-type antigens such as those described above. The analogs may be more soluble or more stable than wild type antigen, and may also contain mutations or modifications rendering the antigen more immunologically active. Also useful in the compositions and 5 methods of the present invention are peptides or proteins which have amino acid sequences homologous with a desired antigen's amino acid sequence, where the homologous antigen induces an immune response to the respective pathogen.

[0071] "Homologous" as used herein refers to the subunit sequence similarity between two 10 polymeric molecules, e.g., between two nucleic acid molecules (e.g., two DNA molecules or two RNA molecules) or two polypeptide molecules. When a subunit position in both molecules is occupied by the same monomeric subunit, e.g., if a position in each of two DNA molecules is occupied by adenine, then they are homologous at that position. The homology between two sequences is a direct function of the number of matching or homologous positions, e.g., if half 15 (e.g. five positions in a polymer ten subunits in length) of the positions in two compound sequences are homologous then the two sequences are 50% homologous, if 90% of the positions, e.g., 9 of 10, are matched or homologous, the two sequences share 90% homology. By way of example, the DNA sequences 5'-CCGTTA-3' and 5'-GCGTAT-3' share 50% homology. By the term "substantially homologous" as used herein, is meant DNA or RNA 20 which is about 50% homologous, more preferably about 70% homologous, even more preferably about 80% homologous and most preferably about 90% homologous to the desired nucleic acid. Genes which are homologous to the desired antigen-encoding sequence should be construed to be included in the invention provided they encode a protein or polypeptide having a biological activity substantially similar to that of the desired antigen. Where in this text, 25 protein and/or DNA sequences are defined by their percent homologies or identities to identified sequences, the algorithms used to calculate the percent homologies or percent identities include the following: the Smith-Waterman algorithm (J. F. Collins *et al*, *Comput. Appl. Biosci.*, (1988) 4:67-72; J. F. Collins *et al*, *Molecular Sequence Comparison and Alignment*, (M. J. Bishop *et al*, eds.) *In Practical Approach Series: Nucleic Acid and Protein Sequence Analysis* 30 XVIII, IRL Press: Oxford, England, UK (1987) 417), and the BLAST and FASTA programs (E. G. Shpaer *et al*, 1996, *Genomics*, 38:179-191). These references are incorporated herein by reference.

[0072] Analogs of the antigens described herein can differ from naturally occurring proteins or 35 peptides by conservative amino acid sequence differences or by modifications which do not affect sequence, or by both. For example, conservative amino acid changes may be made, which although they alter the primary sequence of the protein or peptide, do not normally alter its function. Modifications (which do not normally alter primary sequence) include *in vivo*, or *in vitro* chemical derivatization of polypeptides, e.g., acetylation, or carboxylation. Also

contemplated as antigens are proteins modified by glycosylation, e.g., those made by modifying the glycosylation patterns of a polypeptide during its synthesis and processing or in further processing steps; e.g., by exposing the polypeptide to enzymes which affect glycosylation, e.g., mammalian glycosylating or deglycosylating enzymes. Also contemplated as antigens are 5 amino acid sequences which have phosphorylated amino acid residues, e.g., phosphotyrosine, phosphoserine, or phosphothreonine. Also contemplated as antigens are polypeptides which have been modified using ordinary molecular biological techniques so as to improve their resistance to proteolytic degradation or to optimize solubility properties. Analogs of such polypeptides include those containing residues other than naturally occurring L-amino acids, 10 e.g., D-amino acids or non-naturally occurring synthetic amino acids.

[0073] The antigens of the present invention are not limited to products of any of the specific exemplary processes listed herein. In addition to substantially full length polypeptides, the antigens useful in the present invention include immunologically active fragments of the 15 polypeptides. For example, the antigen may be a fragment of a complete antigen including at least one epitope. "Epitope" as used herein refers to any antigenic determinant on an antigen to which the paratope of an antibody can bind. Epitopic determinants usually consist of chemically active surface groupings of molecules such as, e.g., amino acids or sugar side chains and usually have specific three-dimensional structural characteristics. Particularly 20 preferred embodiments of the compositions and methods of the present invention include combination antigens which include multiple epitopes from the same target antigen, or epitopes from two or more different target antigens (i.e., polytope vaccines). For example, the combination antigens can be the same or different type such as, e.g., a peptide-peptide antigen, glycolipid-peptide antigen, or glycolipid-glycolipid antigen.

25 [0074] A polypeptide or antigen is "immunologically active" if it induces an immune response to a target antigen or pathogen. "Vaccine" as used herein refers to a composition comprising a target antigen that stimulates a specific immune response to that target antigen.

30 [0075] "Adjuvant" as used herein refers to any substance which can stimulate or enhance the stimulation of an immune responses. Some adjuvants can cause activation of a cell of the immune system, for example, an adjuvant can cause an immune cell to produce and secrete cytokines. Examples of adjuvants that can cause activation of a cell of the immune system include, but are not limited to, saponins purified from the bark of the *Q. saponaria* tree, such as 35 QS21 (a glycolipid that elutes in the 21st peak with HPLC fractionation; Aquila Biopharmaceuticals, Inc., Worcester, Mass.); poly(di(carboxylatophenoxy)phosphazene (PCPP polymer; Virus Research Institute, USA); derivatives of lipopolysaccharides such as monophosphoryl lipid A (MPL; Ribi ImmunoChem Research, Inc., Hamilton, Mont.), muramyl dipeptide (MDP; Ribi) and threonyl-muramyl dipeptide (t-MDP; Ribi); OM-174 (a glucosamine

disaccharide related to lipid A; OM Pharma SA, Meyrin, Switzerland); and Leishmania elongation factor (a purified Leishmania protein; Corixa Corporation, Seattle, Wash.).

Traditional adjuvants are well known in the art and include, for example, aluminum phosphate or hydroxide salts ("alum").

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[0076] As compared to known adjuvants, the present invention provides improved adjuvants comprising combinations of lipids and nucleic acids that act synergistically to stimulate enhanced, Th-1 biased immune responses. In preferred embodiments, such compositions of the present invention comprise a nucleic acid component and a lipid component. Preferably the nucleic acid component comprises at least one oligonucleotide, more preferably at least one ODN, and most preferably at least one ODN comprising at least one CpG motif.

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[0077] In preferred embodiments the immunostimulatory compositions used in the methods of the present invention comprise a lipid component comprising a lipid membrane that encapsulates a therapeutic agent. As used herein "liposomal particle," "liposome," "lipid vesicle," and "liposomal vesicle," or grammatical equivalents thereof, may be used interchangeably and refer to structures, particles, complexes, or formulations comprising lipid-containing membranes which enclose or encapsulate an aqueous interior. In preferred embodiments, the liposomes enclose or encapsulate therapeutic agents, e.g., nucleic acids.

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20 The liposomes may have one or more lipid membranes. Liposomes having one lipid-containing membrane are referred to herein as "unilamellar." Liposomes having multiple lipid-containing membranes are referred to herein as "multilamellar." "Lipid bilayer" as used herein refers to a lipid-containing membrane having two layers. In preferred embodiments, the liposomes are multilamellar.

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[0078] The immunostimulatory compositions used in the methods of the present invention generally comprise lipid particles encapsulating at least one methylated nucleic acid.

Nucleic Acids

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[0079] Nucleic acids suitable for use in the compositions of the present invention include, for example, DNA or RNA. Preferably the nucleic acids are oligonucleotides, more preferably ODNs, and most preferably an ODN comprising an ISS ("ISS ODN") and at least one methylated cytosine.

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[0080] "Nucleic acids" as used herein refer to multiple nucleotides (i.e., molecules comprising a sugar (e.g. ribose or deoxyribose) linked to a phosphate group and to an exchangeable organic base, which is either a substituted pyrimidine (e.g. cytosine (C), thymine (T) or uracil (U)) or a substituted purine (e.g. adenine (A) or guanine (G)). Nucleic acids may be, for example DNA or RNA. Preferably the nucleic acids are oligoribonucleotides and more preferably ODNs. Nucleic

acids may also be polynucleosides, i.e., a polynucleotide minus the phosphate and any other organic base containing polymer. The immunostimulatory compositions of the present invention comprise a nucleic acid component. "Nucleic acid component" as used herein with reference to compositions of the present invention refers to a component comprising nucleic acids.

[0081] In a preferred embodiment, the oligonucleotides are single stranded and in the range of 5 - 50 nucleotides ("nt") in length. However, any oligonucleotides may be used including, for example, large double stranded plasmid DNA in the range of 500 - 50,000 base pairs ("bp").

[0082] Nucleic acids useful in the compositions and methods of the present invention can be obtained from known sources or isolated using methods well known in the art. The nucleic acids can also be prepared by recombinant or synthetic methods which are equally well known in the art. Such nucleic acids can then be encapsulated in lipid particles and the resulting compositions tested for immunostimulatory activity using the methods of the present invention as described herein.

[0083] For use *in vivo*, nucleic acids may be resistant to degradation (e.g., via endo-and exo-nucleases). Secondary structures, such as stem loops, can stabilize nucleic acids against degradation. Alternatively, nucleic acid stabilization can be accomplished via phosphate backbone modifications. A preferred stabilized nucleic acid has at least a partial phosphorothioate modified backbone. Phosphorothioates may be synthesized using automated techniques employing either phosphoramidate or H-phosphonate chemistries. Aryl-and alkyl-phosphonates can be made, e.g., as described in U.S. Patent No. 4,469,863; and alkylphosphotriesters (in which the charged oxygen moiety is alkylated as described in U.S. Patent No. 5,023,243 and European Patent No. 092,574) can be prepared by automated solid phase synthesis using commercially available reagents. Methods for making other DNA backbone modifications and substitutions have been described (Uhlmann and Peyman, *Chem. Rev.* 90:544, 1990; Goodchild, *Bioconjugate Chem.* 1:165, 1990). As described herein, however, the methods and compositions of the present invention alleviate the need to include such modifications to the subject nucleic acids.

[0084] Thus, oligonucleotides useful in the compositions and methods of the present invention may have a modified phosphate backbone such as, e.g., phosphorothioate, methylphosphonate, methylphosphorothioate, phosphorodithioate, and combinations thereof with each other and/or with phosphodiester oligonucleotide. In addition, other modified oligonucleotides include: nonionic DNA analogs, such as alkyl- and aryl-phosphates (in which the charged phosphonate oxygen is replaced by an alkyl or aryl group), phosphodiester and alkylphosphotriesters, in which the charged oxygen moiety is alkylated. As demonstrated

herein, PO ODN may be preferred where cellular immune responses are desired, while modified ODN such as, e.g., PS ODN may be preferred where humoral responses are desired.

[0085] Numerous other chemical modifications to the base, sugar or linkage moieties are also 5 useful. Bases may be methylated or unmethylated. In the preferred embodiments, methyl or hydroxymethyl groups are attached to the carbon-4 position (4-mC) or carbon-5 position (5-mC) of at least one cytosine. The methylated cytosine is preferably located within a CpG motif in the nucleic acid sequence. Alternatively or additionally, the sugar moiety may be modified with a methyl group as described in the art.

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[0086] Nucleic acid sequences useful in the compositions and methods of the present invention may be complementary to patient/subject mRNA, such as antisense oligonucleotides, or they may be foreign or non-complementary (e.g., the nucleotide sequences do not specifically hybridize to the patient/subject genome). The nucleotide sequences may be expressed and the 15 resulting expression products may be RNA and/or protein. In addition, such nucleotide sequences may be linked to appropriate promoters and expression elements, and may be contained in an expression vector. Nucleotide sequences useful in the composition and methods of the present invention may be ISS, such as certain palindromes leading to hairpin secondary structures (see Yamamoto S., *et al.* (1992) *J. Immunol.* 148: 4072-4076), or CpG 20 motifs, or other known ISS features (such as multi-G domains, see WO 96/11266). In a particularly preferred embodiment, the nucleotide sequence comprises at least one CpG motif having a methylated cytosine.

[0087] The nucleic acids of the present invention can be synthesized *de novo* using any of a 25 number of procedures well known in the art. For example, the *b*-cyanoethyl phosphoramidite method (Beaucage, S. L., and Caruthers, M. H., *Tet. Let.* 22:1859, 1981); nucleoside H-phosphonate method (Garegg *et al.*, *Tet. Let.* 27:4051-4054, 1986; Froehler *et al.*, *Nucl. Acid. Res.* 14:5399-5407, 1986; Garegg *et al.*, *Tet. Let.* 27:4055-4058, 1986, Gaffney *et al.*, *Tet. Let.* 29:2619-2622, 1988). These chemistries can be performed by a variety of automated 30 oligonucleotide synthesizers available in the market. Also, CpG dinucleotides can be produced on a large scale in plasmids, (see Sambrook, T., *et al.*, *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor laboratory Press, New York, 1989). Such plasmids may also encode other genes to be expressed such as an antigen-encoding gene in the case of a DNA vaccine. Oligonucleotides can be prepared from existing nucleic acid sequences (e.g., 35 genomic or cDNA) using known techniques, such as those employing restriction enzymes, exonucleases or endonucleases.

[0088] For administration *in vivo*, compositions of the present invention, including components of the compositions, e.g., a lipid component or a nucleic acid component, may be associated

with a molecule that results in higher affinity binding to target cell (e.g., B-cell, monocytic cell and natural killer (NK) cell) surfaces and/or increased cellular uptake by target cells. The compositions of the present invention, including components of the compositions, can be ionically or covalently associated with desired molecules using techniques which are well known 5 in the art. A variety of coupling or cross-linking agents can be used, e.g., protein A, carbodiimide, and N-succinimidyl-3-(2-pyridylthio) propionate (SPDP).

[0089] The immune stimulating activity of a nucleic acid sequence in an organism can be determined by simple experimentation, for example, by comparing the sequence in question 10 with other immunostimulatory agents, e.g., other adjuvants, or ISS; or by detecting or measuring the immunostimulatory activity of the sequence in question, e.g., by detecting or measuring the activation of host defense mechanisms or the activation of immune system components. Such assays are well known in the art. Also, one of skill in the art would know how to identify the optimal oligonucleotides useful for a particular mammalian species of 15 interest using routine assays described herein and/or known in the art.

[0090] Specific nucleic acid sequences of ODNs suitable for use in the compositions and methods of the invention are described in U.S. Patent Appln. 60/379,343, U.S. Patent Appln. No. 09/649,527, Int. Publ. WO 02/069369, Int. Publ. No. WO 01/15726, U.S. Patent No. 20 6,406,705, and Raney *et al.*, Journal of Pharmacology and Experimental Therapeutics, 298:1185-1192 (2001), which are all incorporated herein by reference. Exemplary sequences of the ODNs include, but are not limited to, those nucleic acid sequences shown in Table 1. In preferred embodiments, ODNs used in the compositions and methods of the present invention have a phosphodiester ("PO") backbone or a phosphorothioate ("PS") backbone, and at least 25 one methylated cytosine residue in the CpG motif.

Table 1

ODN NAME	ODN SEQ ID NO	ODN SEQUENCE (5'-3')
ODN 1 (INX-6295) human c-myc	SEQ ID NO: 2	5'-TAACGTTGAGGGGCAT-3
* ODN 1m (INX-6303)	SEQ ID NO: 4	5'-TAAZGTTGAGGGGCAT-3
ODN 2 (INX-1826)	SEQ ID NO: 1	5'-TCCATGACGTTCCCTGACGTT-3
* ODN 2m (INX-1826m)	SEQ ID NO: 31	5'-TCCATGAZGTTCCCTGAZGTT-3
ODN 3 (INX-6300)	SEQ ID NO: 3	5'-TAAGCATAACGGGGTGT-3
ODN 5 (INX-5001)	SEQ ID NO: 5	5'-AACGTT-3

ODN 6 (INX-3002)	SEQ ID NO: 6	5'-GATGCTGTGTCGGGGTCTCCGGGC-3'
ODN 7 (INX-2006)	SEQ ID NO: 7	5'-TCGTCGTTTGTCTGGTT-3'
ODN 7m (INX-2006m)	SEQ ID NO: 7	5'-TZGTZGTTTGTZGTTTGTZGTT-3'
ODN 8 (INX-1982)	SEQ ID NO: 8	5'-TCCAGGACTCTCTCAGGTT-3'
ODN 9 (INX-G3139)	SEQ ID NO: 9	5'-TCTCCCAGCGTGCGCCAT-3'
ODN 10 (PS-3082) murine Intracellular Adhesion Molecule-1	SEQ ID NO: 10	5'-TGCAATCCCCAGGCCACCAT-3
ODN 11 (PS-2302) human Intracellular Adhesion Molecule-1	SEQ ID NO: 11	5'-GCCCAAGCTGGCATCCGTCA-3'
ODN 12 (PS-8997) human Intracellular Adhesion Molecule-1	SEQ ID NO: 12	5'-GCCCAAGCTGGCATCCGTCA-3'
ODN 13 (US3) human erb-B-2	SEQ ID NO: 13	5'-GGT GCTCACTGC GGC-3'
ODN 14 (LR-3280) human c-myc	SEQ ID NO: 14	5'-AACCGTTGAGGGCAT-3'
ODN 15 (LR-3001) human c-myc	SEQ ID NO: 15	5'-TAT GCT GTG CCG GGG TCT TCG GGC-3'
ODN 16 (Inx-6298)	SEQ ID NO: 16	5'-GTGCCGGGGTCTCGGGC-3'
ODN 17 (hIGF-1R) human Insulin Growth Factor 1 – Receptor	SEQ ID NO: 17	5'-GGACCCCTCCTCCGGAGCC-3'
ODN 18 (LR-52) human Insulin Growth Factor 1 – Receptor	SEQ ID NO: 18	5'-TCC TCC GGA GCC AGA CTT-3'
ODN 19 (hEGFR) human Epidermal Growth Factor – Receptor	SEQ ID NO: 19	5'-AAC GTT GAG GGG CAT-3'
ODN 20 (EGFR) Epidermal Growth Factor – Receptor	SEQ ID NO: 20	5'-CCGTGGTCA TGCTCC-3'
ODN 21 (hVEGF) human Vascular Endothelial Growth Factor	SEQ ID NO: 21	5'-CAG CCTGGCTACCG CCTTGG-3'

ODN 22 (PS-4189) murine Phosphokinase C – alpha	SEQ ID NO: 22	5'-CAG CCA TGG TTC CCC CCA AC-3'
ODN 23 (PS-3521)	SEQ ID NO: 23	5'-GTT CTC GCT GGT GAG TTT CA-3'
ODN 24 (hBcl-2) human Bcl-2	SEQ ID NO: 24	5'-TCT CCCAGCGTGCGCCAT-3'
ODN 25 (hC-Raf-1) human C-Raf-s	SEQ ID NO: 25	5'-GTG CTC CAT TGA TGC-3'
ODN #26 (hVEGF-R1) human Vascular Endothelial Growth Factor Receptor-1	SEQ ID NO: 26	5'-GAGUUCUGAUGAGGCCGAAAGGCCG AAAGUCUG-3'
ODN #27	SEQ ID NO: 27	5'-RRCGYY-3'
ODN #28 (INX-3280)	SEQ ID NO: 28	5'-AACGTTGAGGGGCAT-3'
ODN #29 (INX-6302)	SEQ ID NO: 29	5'-CAACGTTATGGGGAGA-3'
ODN #30 (INX-6298) human c-myc	SEQ ID NO: 30	5'-TAACGTTGAGGGGCAT-3'

- “Z” represents a methylated cytosine residue.
- Note: ODN 14 is a 15-mer oligonucleotide and ODN 1 is the same oligonucleotide having a thymidine added onto the 5' end making ODN 1 into a 16-mer. No difference in biological activity between ODN 14 and ODN 1 has been detected and both exhibit similar immunostimulatory activity (Mui *et al.*, 2001)

Lipids and other components

[0091] Lipid formulations and methods of preparing liposomes as delivery vehicles are known in the art, and any of number of such formulations may find advantageous use herein, including those described in U.S. Patent No. 6,465,439, U.S. Patent No. 6,379,698, U.S. Patent No. 6,365,611, and U.S. Patent No. 6,093,816, the disclosures of which are incorporated herein by reference. Preferred lipid formulations are the lipid particle formulations described herein and more fully described in, for example, U.S. Patent No. 5,785,992, U.S. Patent No. 6,287,591, U.S. Patent No. 6,287,591 B1, co-pending U.S. Patent Appln. Ser. No. 60/379,343, and co-pending U.S. Patent Appln. Ser. No. 09/649,527 all incorporated herein by reference.

[0092] In one preferred embodiment, the preferred lipid formulation is DSPC, DODMA, Chol, and PEG-DMG having a ratio of 20:25:45:10 mol/mol. As used herein, the molar amount of each lipid in a lipid formulation is given in the same order that the lipid is listed (e.g., the ratio of 20 DSPC to 25 DODMA to 45 Chol to 10 PEG-DMG or “20:25:45:10”). In alternate embodiments the DSPC may be replaced with POPC, the DODMA

replaced with DODAP and the PEG-DMG replaced with PEGCer14 or PEGCer20.

[0093] The term "lipid" refers to a group of organic compounds that are esters of fatty acids and are characterized by being insoluble in water but soluble in many organic solvents. They are 5 usually divided in at least three classes: (1) "simple lipids" which include fats and oils as well as waxes; (2) "compound lipids" which include phospholipids and glycolipids; and (3) "derived lipids" such as steroids and compounds derived from lipid manipulations. A wide variety of lipids may be used with the invention, some of which are described below.

10 [0094] The term "charged lipid" refers to a lipid species having either a cationic charge or negative charge or which is a zwitterion which is not net neutrally charged, and generally requires reference to the pH of the solution in which the lipid is found.

[0095] Cationic charged lipids at physiological pH include, but are not limited to, N,N-dioleyl-15 N,N-dimethylammonium chloride ("DODAC"); N-(2,3-dioleyloxy)propyl)-N,N,N-trimethylammonium chloride ("DOTMA"); N,N-distearyl-N,N-dimethylammonium bromide ("DDAB"); N-(2,3-dioleyloxy)propyl)-N,N,N-trimethylammonium chloride ("DOTAP"); 3b-(N-(N',N'-dimethylaminoethane)-carbamoyl)cholesterol ("DC-Chol") and N-(1,2-dimyristyloxyprop-3-yl)-N,N-dimethyl-N-hydroxyethyl ammonium bromide ("DMRIE"). Additionally, a number of 20 commercial preparations of cationic lipids are available which can be used in the present invention. These include, for example, Lipofectin™ (commercially available cationic liposomes comprising DOTMA and 1,2-dioleoyl-sn-3-phosphoethanolamine ("DOPE"), from GIBCO/BRL, Grand Island, New York, U.S.A); and Lipofectamine™ (commercially available cationic liposomes comprising N-(1-(2,3-dioleyloxy)propyl)-N-(2-(sperminecarboxamido)ethyl)-N,N-25 dimethylammonium trifluoroacetate ("DOSPA").

[0096] Some cationic charged lipids are titratable, that is to say they have a pKa at or near physiological pH, with the significant consequence for this invention that they are strongly cationic in mild acid conditions and weakly (or not) cationic at physiological pH. Such cationic 30 charged lipids include, but are not limited to, N-(2,3-dioleyloxy)propyl)-N,N-dimethylammonium chloride ("DODMA") and 1,2-Dioleoyl-3-dimethylammonium-propane ("DODAP"). DMDMA is also a useful titratable cationic lipid.

[0097] Anionic charged lipids at physiological pH include, but are not limited to, phosphatidyl 35 inositol, phosphatidyl serine, phosphatidyl glycerol, phosphatidic acid, diphosphatidyl glycerol, poly(ethylene glycol)-phosphatidyl ethanolamine, dimyristoylphosphatidyl glycerol, dioleoylphosphatidyl glycerol, dilauryloylphosphatidyl glycerol, dipalmitoylphosphatidyl glycerol, distearoylphosphatidyl glycerol, dimyristoyl phosphatic acid, dipalmitoyl phosphatic acid, dimyristoyl phosphatidyl serine, dipalmitoyl phosphatidyl serine, brain phosphatidyl serine, and

the like.

[0098] Some anionic charged lipids may be titratable, that is to say they would have a pKa at or near physiological pH, with the significant consequence for this invention that they are 5 strongly anionic in mild base conditions and weakly (or not) anionic at physiological pH. Such anionic charged lipids can be identified by one skilled in the art based on the principles disclosed herein.

[0099] The term "neutral lipid" refers to any of a number of lipid species which exist either in an 10 uncharged or neutral zwitterionic form at physiological pH. Such lipids include, for example, diacylphosphatidylcholine, diacylphosphatidylethanolamine, ceramide, sphingomyelin, cephalin, cholesterol, cerebrosides and diacylglycerols.

[00100] Certain preferred lipid formulations used in the invention include aggregation 15 preventing compounds such as PEG-lipids or polyamide oligomer-lipids (such as an ATTA-lipid), and other steric-barrier or "stealth"-lipids, detergents, and the like. Such lipids are described in U.S. Patent No. 4,320,121, U.S. Patent No. 5,820,873, U.S. Patent No. 5,885,613, Int. Publ. No. WO 98/51278, and U.S. Pat. Appln. Serial No. 09/218,988 relating to polyamide oligomers, all incorporated herein by reference. These lipids and detergent compounds prevent 20 precipitation and aggregation of formulations containing oppositely charged lipids and therapeutic agents. These lipids may also be employed to improve circulation lifetime *in vivo* (see Klibanov *et al.* (1990) FEBS Letters, 268 (1): 235-237), or they may be selected to rapidly exchange out of the formulation *in vivo* (see U.S. Patent No. 5,885,613, incorporated herein by reference).

25 [00101] A preferred embodiment of the invention employs exchangeable steric-barrier lipids (as described in U.S. Patent No. 5,820,873, U.S. Patent No. 5,885,613, and U.S. Pat. Appln. Ser. No. 09/094540 and U.S. Pat. No. 6,320,017, all incorporated herein by reference). Exchangeable steric-barrier lipids such as PEG2000-CerC14 and ATTA8-CerC14 are steric- 30 barrier lipids which rapidly exchange out of the outer monolayer of a lipid particle upon administration to a subject/patient. Each such lipid has a characteristic rate at which it will exchange out of a particle depending on a variety of factors including acyl chain length, saturation, size of steric barrier moiety, membrane composition and serum composition, etc. Such lipids are useful in preventing aggregation during particle formation, and their accelerated 35 departure from the particle upon administration provides benefits, such as programmable fusogenicity and particle destabilizing activity, as described in the above noted patent submissions.

[00102] Some lipid particle formulations may employ targeting moieties designed to encourage localization of liposomes at certain target cells or target tissues. Targeting moieties may be associated with the outer bilayer of the lipid particle (i.e., by direct conjugation, hydrophobic interaction or otherwise) during formulation or post-formulation. These methods are well known in the art. In addition, some lipid particle formulations may employ fusogenic polymers such as PEAA, hemagglutinin, other lipo-peptides (see U.S. Pat. No. 6,417,326, and U.S. Pat. Appln. Ser. No. 09/674,191, all incorporated herein by reference) and other features useful for *in vivo* and/or intracellular delivery.

5 [00103] In another preferred embodiment, the lipid component lipid particles of the present invention comprises sphingomyelin and cholesterol ("sphingosomes"). In a preferred embodiment, the lipid particles used in the compositions and methods of the present invention are comprised of sphingomyelin and cholesterol and have an acidic intraliposomal pH. The lipid particles comprising sphingomyelin and cholesterol have several advantages when compared to other formulations. The sphingomyelin/cholesterol combination produces liposomes which have extended circulation lifetimes, are much more stable to acid hydrolysis, have significantly better drug retention characteristics, have better loading characteristics into tumors and the like, and show significantly better anti-tumor efficacy than other liposomal formulations tested.

10 [00104] In a preferred embodiment, the lipid particles of the present invention comprise a cationic compound of Formula I and at least one neutral lipid as follows (and fully described in U.S. Pat. Serial No. 5,785,992, incorporated herein by reference). In a preferred embodiment, the LNA formulations of the present invention comprise a cationic compound of Formula I and at least one neutral lipid as follows (and fully described in U.S. Pat. Serial No. 5,785,992, incorporated herein by reference).

15 [00105] In Formula I, R¹ and R² are each independently C₁ to C₃ alkyl. Y and Z are alkyl or alkenyl chains and are each independently: --CH₂CH₂CH₂CH₂CH₂--, --CH=CHCH₂CH₂CH₂--, --CH₂CH=CHCH₂CH₂--, --CH₂CH=CHCH=CH₂--, --CH=CHCH=CHCH₂--, --CH=CHCH=CHCH=CH₂--, --CH=CHCH=CHCH=CH=CH₂--, --CH=CHCH=CHCH=CH=CH=CH₂--, --CH=CHCH=CHCH=CH=CH=CH=CH₂--, or --CH₂CH=CHCH=CHCH=CH=CH₂--, with the proviso that Y and Z are not both --CH₂CH₂CH₂CH₂CH₂--. The letters n and q denote integers of from 3 to 7, while the letters m and p denote integers of from 4 to 9, with the proviso that the sums n+m and q+p are each integers of from 10 to 14. The symbol X' represents a pharmaceutically



acceptable anion. In the above formula, the orientation of the double bond can be either cis or trans, however the cis isomers are generally preferred.

5 [00106] In another preferred embodiment, the cationic compounds are of Formula I, wherein R¹ and R² are methyl and Y and Z are each independently: --CH=CHCH₂CH₂CH₂--, --CH₂CH=CHCH₂CH₂--, --CH₂CH₂CH=CHCH₂-- or --CH₂CH₂CH₂CH=CH--. In preferred embodiments, R¹ and R² are methyl; Y and Z are each --CH=CHCH₂CH₂CH₂--; n and q are both 7; and m and p are both 5. In another preferred embodiment, the cationic compound is 10 DODAC (N,N-dioleyl-N,N-dimethylammonium chloride). DODAC is a known in the art and is a compound used extensively as an additive in detergents and shampoos. DODA is also used as a co-lipid in liposomal compositions with other detergents (see, Takahashi, *et al.*, GB 2147243).

15 [00107] The neutral lipids in the LNA formulations of the present invention can be any of a variety of neutral lipids which are typically used in detergents, or for the formation of micelles or liposomes. Examples of neutral lipids which are useful in the present compositions are, but are not limited to, diacylphosphatidylcholine, diacylphosphatidylethanolamine, ceramide, sphingomyelin, cephalin, cardiolipin, and cerebrosides. In a preferred embodiment, the present compositions will include one or more neutral lipids which are diacylphosphatidylcholine, diacylphosphatidylethanolamine, ceramide or sphingomyelin. The 20 acyl groups in these neutral lipids are preferably acyl groups derived from fatty acids having C₁₀-C₂₄ carbon chains. More preferably the acyl groups are lauroyl, myristoyl, palmitoyl, stearoyl or oleoyl. In particularly preferred embodiments, the neutral lipid will be 1,2-sn-dioleoylphosphatidylethanolamine.

25 [00108] The anion, X⁻, can similarly be any of a variety a pharmaceutically acceptable anions. These anions can be organic or inorganic, including for example, Br⁻, Cl⁻, F⁻, I⁻, sulfate, phosphate, acetate, nitrate, benzoate, citrate, glutamate, and lactate. In preferred embodiments, X⁻ is Cl⁻ or AcO⁻.

30 [00109] In addition to the other components described herein, the compositions of the present invention may contain a pharmaceutically acceptable carrier. Pharmaceutically acceptable carriers are well-known in the art. The choice of carrier is determined in part by the particular composition to be administered as well as by the particular method used to administer the composition. Preferably, the pharmaceutical carrier is in solution, in water or saline.

35 [00110] In the compositions of the present invention, the ratio of cationic compound to neutral lipid is preferably within a range of from about 25:75 (cationic compound:neutral lipid), or

preferably to 75:25 (cationic compound:neutral lipid), or preferably about 50:50.

[00111] The cationic compounds which are used in the compositions of the present invention can be prepared by methods known to those of skill in the art using standard synthetic reactions

5 (see March, Advanced Organic Chemistry, 4th Ed., Wiley-Interscience, NY, N.Y. (1992), incorporated herein by reference). For example, the synthesis of OSDAC can be carried out by first treating oleylamine with formaldehyde and sodium cyanoborohydride under conditions which result in the reductive alkylation of the amine. This approach provides dimethyl oleylamine, which can then be alkylated with stearyl bromide to form the corresponding 10 ammonium salt. Anion exchange results in the formation of OSDAC. Dimethyloleylamine can also be synthesized by treatment of oleyl bromide with a large excess of dimethylamine, and further derivatized as described above.

[00112] For cationic compounds in which both fatty acid chains are unsaturated (i.e., DODAC),

15 the following general procedure can be used. An unsaturated acid (i.e., oleic acid) can be converted to its corresponding acyl chloride with such reagents as oxalyl chloride, thionyl chloride, PCl_3 or PCl_5 . The acyl chloride can be treated with an unsaturated amine (i.e., oleylamine) to provide the corresponding amide. Reduction of the amide with, for example, lithium aluminum hydride provides a secondary amine wherein both alkyl groups are 20 unsaturated long chain alkyl groups. The secondary amine can then be treated with alkyl halides such as methyl iodide to provide a quaternary ammonium compound. Anion exchange can then be carried out to provide cationic compounds having the desired pharmaceutically acceptable anion. The alkylamine precursor can be synthesized in a similar manner. For example, treatment of an alkyl halide with a methanolic solution of ammonia in large excess will 25 produce the required amine after purification. Alternatively, an acyl chloride, produced by treatment of the appropriate carboxylic acid with oxalyl chloride, can be reacted with ammonia to produce an amide. Reduction of the amide with LiAlH_4 will provide the required alkylamine.

[00113] In preferred embodiments, the pharmaceutical compositions of the present invention

30 are formulated as micelles or liposomes. Micelles containing the cationic compounds and neutral lipids of the present invention can be prepared by methods well known in the art. In addition to the micellar formulations of the present compositions, the present invention also provides micellar formulations which include other species such as lysophosphatidylcholine, lysophosphatidylethanolamine, lysophosphatidylserine, lysophosphatidylglycerol, 35 phosphatidylethanolamine-polyoxyethylene conjugate, ceramide-polyoxyethylene conjugate or phosphatidic acid-polyoxyethylene conjugate.

[00114] The polyoxyethylene conjugates which are used in the compositions of the present invention can be prepared by combining the conjugating group (i.e. phosphatidic acid or

phosphatidylethanolamine) with an appropriately functionalized polyoxyethylene derivative. For example, phosphatidylethanolamine can be combined with omega-methoxypolyethyleneglycol succinate to provide a phosphatidylethanolamine-polyoxyethylene conjugate (see, e.g., Parr, *et al.*, *Biochim. Biophys. Acta* 1195:21-30 (1994), incorporated herein by reference).

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[00115] The selection of neutral lipids for use in the compositions and methods of the present invention is generally guided by consideration of, e.g., liposome size and stability of the liposomes in the bloodstream. As described above, the neutral lipid component in the liposomes is a lipid having two acyl groups, (i.e., diacylphosphatidylcholine and

10 diacylphosphatidyl-ethanolamine). Lipids having a variety of acyl chain groups of varying chain length and degree of saturation are available or may be isolated or synthesized by well-known techniques. In general, less saturated lipids are more easily sized, particularly when the liposomes must be sized below about 0.3 microns, for purposes of filter sterilization. In one group of embodiments, lipids containing saturated fatty acids with carbon chain lengths in the 15 range of C14 to C22 are preferred. In another group of embodiments, lipids with mono or diunsaturated fatty acids with carbon chain lengths in the range of C14 to C22 are used. Additionally, lipids having mixtures of saturated and unsaturated fatty acid chains can be used.

[00116] Liposomes useful in the compositions and methods of the present invention may also 20 be composed of sphingomyelin or phospholipids with other head groups, such as serine and inositol. Still other liposomes useful in the present invention will include cholesterol, diglycerides, ceramides, phosphatidylethanolamine-polyoxyethylene conjugates, phosphatidic acid-polyoxyethylene conjugates, or polyethylene glycol-ceramide conjugates (e.g., PEG-Cer-C14 or PEG-Cer-C20). Methods used in sizing and filter-sterilizing liposomes are discussed 25 below.

[00117] A variety of methods are known in the art for preparing liposomes (see e.g., Szoka *et al.*, *Ann. Rev. Biophys. Bioeng.* 9:467 (1980), U.S. Pat. Nos. 4,235,871, 4,501,728, 4,837,028, the text *Liposomes*, Marc J. Ostro, ed., Marcel Dekker, Inc., New York, 1983, Chapter 1, and 30 Hope, *et al.*, *Chem. Phys. Lip.* 40:89 (1986), all of which are incorporated herein by reference). One known method produces multilamellar vesicles of heterogeneous sizes. In this method, the vesicle-forming lipids are dissolved in a suitable organic solvent or solvent system and dried under vacuum or an inert gas to form a thin lipid film. If desired, the film may be redissolved in a suitable solvent, such as tertiary butanol, and then lyophilized to form a more homogeneous 35 lipid mixture which is in a more easily hydrated powder-like form. This film is covered with an aqueous buffered solution and allowed to hydrate, typically over a 15-60 minute period with agitation. The size distribution of the resulting multilamellar vesicles can be shifted toward smaller sizes by hydrating the lipids under more vigorous agitation conditions or by adding

solubilizing detergents such as deoxycholate.

[00118] Following liposome preparation, the liposomes may be sized to achieve a desired size range and relatively narrow distribution of liposome sizes. A size range of about 0.2-0.4 microns allows the liposome suspension to be sterilized by filtration through a conventional filter, typically a 0.22 micron filter. The filter sterilization method can be carried out on a high through-put basis if the liposomes have been sized down to about 0.2-0.4 microns.

[00119] Several techniques are available for sizing liposomes to a desired size. One sizing method is described in U.S. Patent No. 4,737,323, incorporated herein by reference. 10 Sonicating a liposome suspension either by bath or probe sonication produces a progressive size reduction down to small unilamellar vesicles less than about 0.05 microns in size. Homogenization is another method which relies on shearing energy to fragment large 15 liposomes into smaller ones. In a typical homogenization procedure, multilamellar vesicles are recirculated through a standard emulsion homogenizer until selected liposome sizes, typically between about 0.1 and 0.5 microns, are observed. In both methods, the particle size distribution can be monitored by conventional laser-beam particle size discrimination.

[00120] Extrusion of liposomes through a small-pore polycarbonate membrane or an 20 asymmetric ceramic membrane is also an effective method for reducing liposome sizes to a relatively well-defined size distribution. Typically, the suspension is cycled through the membrane one or more times until the desired liposome size distribution is achieved. The liposomes may be extruded through successively smaller-pore membranes, to achieve a gradual reduction in liposome size. For use in the present inventions, liposomes having a size 25 of from about 0.05 microns to about 0.15 microns are preferred.

[00121] As further described below, the compositions of the present invention can be 30 administered to a subject by any known route of administration. Once adsorbed by cells, the liposomes (including the complexes previously described) can be endocytosed by a portion of the cells, exchange lipids with cell membranes, or fuse with the cells. Transfer or incorporation of the polyanionic portion of the complex can take place via any one of these pathways. In particular, when fusion takes place, the liposomal membrane can be integrated into the cell membrane and the contents of the liposome can combine with the intracellular fluid.

35 [00122] As described below in detail, additional components, which may also be therapeutic compounds, may be added to the lipid particles of the present invention to target them to specific cell types. For example, the liposomes can be conjugated to monoclonal antibodies or binding fragments thereof that bind to epitopes present only on specific cell types, such as cancer-related antigens, providing a means for targeting the liposomes following systemic

administration. Alternatively, ligands that bind surface receptors of the target cell types may also be bound to the liposomes. Other means for targeting liposomes may also be employed in the present invention.

5 [00123] Following a separation step as may be necessary to remove free drug from the medium containing the liposome, the liposome suspension is brought to a desired concentration in a pharmaceutically acceptable carrier for administration to the patient or host cells. Many pharmaceutically acceptable carriers may be employed in the compositions and methods of the present invention. A variety of aqueous carriers may be used, e.g., water, 10 buffered water, 0.4% saline, 0.3% glycine, and the like, and may include glycoproteins for enhanced stability, such as albumin, lipoprotein, globulin. Generally, normal buffered saline (135-150 mM NaCl) will be employed as the pharmaceutically acceptable carrier, but other suitable carriers will suffice. These compositions may be sterilized by conventional liposomal sterilization techniques, such as filtration. The compositions may contain pharmaceutically 15 acceptable auxiliary substances as required to approximate physiological conditions, such as pH adjusting and buffering agents, tonicity adjusting agents and the like, for example, sodium acetate, sodium lactate, sodium chloride, potassium chloride, calcium chloride. These compositions may be sterilized techniques referred to above or produced under sterile conditions. The resulting aqueous solutions may be packaged for use or filtered under aseptic 20 conditions and lyophilized, the lyophilized preparation being combined with a sterile aqueous solution prior to administration.

[00124] The concentration of liposomes in the carrier may vary. In preferred embodiments, the concentration of liposomes is about 0.1-200 mg/ml. Persons of skill would know how to vary 25 these concentrations to optimize treatment with different liposome components or for particular patients. For example, the concentration may be increased to lower the fluid load associated with treatment.

[00125] The cells of a subject are usually exposed to the compositions of the present invention 30 by in vivo or ex vivo administration. In the preferred embodiments described herein, the compositions of the present invention are administered systemically, e.g., intravenously, with intramuscular, subcutaneous and topical administration also contemplated. Alternatively, intranasal or intratracheal administration may be used. Intratracheal administration may be provided as a liquid, preferably as an aerosol. For example, nebulizers may be used to create 35 aerosols of droplets of between 70-100 μ m in diameter. It will be understood that droplet size should generally be of greater size than the liposomes.

[00126] Multiple administrations to a patient are contemplated. The dosage schedule of the treatments will be determined by the disease and the patient's condition. Standard treatments

with therapeutic compounds, including immunostimulatory compositions (e.g., vaccines), that are well known in the art may serve as a guide to treatment with liposomes containing the therapeutic compounds. The duration and schedule of treatments may be varied by methods well known to those of skill, but the increased circulation time and decreased in liposome

5 leakage will generally allow the dosages to be adjusted downward from those previously employed. The dose of liposomes of the present invention may vary depending on the clinical condition and size of the animal or patient receiving treatment. The standard dose of the therapeutic compound when not encapsulated may serve as a guide to the dose of the liposome-encapsulated compound. The dose will typically be constant over the course of

10 treatment, although in some cases the dose may vary. Standard physiological parameters may be assessed during treatment that may be used to alter the dose of the liposomes of the invention.

Antigens

15 [00127] As described herein, the liposomal encapsulated methylated nucleic acids may be associated with at least one target antigen. Antigens useful in the compositions and methods of the present invention may be inherently immunogenic, or non-immunogenic, or slightly immunogenic. Examples of antigens include, but are not limited to, synthetic, recombinant, foreign, or homologous antigens. Further examples of antigens include, but are not limited to,

20 HBA – hepatitis B antigen (recombinant or otherwise); other hepatitis peptides; HIV proteins GP120 and GP160; Mycoplasma cell wall lipids; any tumor associated antigen; Carcinoembryonic Antigen (CEA); other embryonic peptides expressed as tumor specific antigens; bacterial cell wall glycolipids; Gangliosides (GM2, GM3); Mycobacterium glycolipids; PGL-1; Ag85B; TBGL; Gonococci lip-oligosaccharide epitope 2C7 from Neisseria gonorrhoeae;

25 Lewis(y); and Globo-H; Tn; TF; STn; PorA; TspA or Viral glycolipids/glycoproteins and surface proteins.

30 [00128] The antigen may be in the form of a peptide antigen or it may be a nucleic acid encoding an antigenic peptide in a form suitable for expression in a subject and presentation to the immune system of the immunized subject. The antigen may also be a glycolipid or a glycopeptide. Further, the antigen may be a complete antigen, or it may be a fragment of a complete antigen including at least one therapeutically relevant epitope. "Combination antigens" as herein refer to antigens having multiple epitopes from the same target antigen, or multiple epitopes from two or more different target antigens (polytope vaccines) originating from

35 the same type of target antigens (e.g., both antigens are peptides or both antigens are glycolipids), or different types of target antigens (e.g., glycolipid antigen and peptide antigen).

[00129] Antigens may be used in the compositions and methods of the present invention in a crude, purified, synthetic, isolated, or recombinant form. Polypeptide or peptide antigens,

(including, for example, antigens that are peptide mimics of polysaccharides) encoded by nucleic acids may also be used in the compositions and methods of the present invention. The term antigen broadly includes any type of molecule which is recognized by a host immune system as being foreign. Antigens include but are not limited to cancer antigens, microbial 5 antigens, and allergens.

[00130] A wide variety of antigens are suitable for use in the formulations of the present invention. Generally, an antigen is a material that is administered to a vertebrate host to immunize the host against the same material. Typically, an antigen comprises material 10 associated with a disease state, such as viral infection, bacterial infection, and various malignancies. These materials may include but are not limited to proteins, peptides, polypeptides, lipids, glycolipids, carbohydrates and DNA.

[00131] The antigen of the lipid formulation may be encapsulated, associated, or mixed with the 15 liposome or lipid particle. In certain embodiments of the present invention, the antigen is encapsulated in the liposome or lipid particle. In other embodiments, the antigen is mixed with the liposome or lipid particle. In other embodiments, the antigen is associated with the liposome or lipid particle. In one aspect, the antigen is adsorbed to the liposome or lipid particle. In other aspects, the antigen is covalently attached to the liposome or lipid particle. 20 Methods used to covalently attach the antigen to the liposome or lipid particle are those standard methods known to those of skill in the art.

[00132] Examples of antigens suitable for use in the present invention include, but are not limited to, polypeptide antigens and DNA antigens. Specific examples of antigens are Hepatitis 25 A, Hepatitis B, small pox, polio, anthrax, influenza, typhus, tetanus, measles, rotavirus, diphtheria, pertussis, tuberculosis, and rubella antigens. In a preferred embodiment, the antigen is a Hepatitis B recombinant antigen. In other aspects, the antigen is a Hepatitis A recombinant antigen. In another aspect, the antigen is a tumor antigen. Examples of such tumor-associated antigens are MUC-1, EBV antigen and antigens associated with Burkitt's 30 lymphoma. In a further aspect, the antigen is a tyrosinase-related protein tumor antigen recombinant antigen. Those of skill in the art will know of other antigens suitable for use in the present invention.

[00133] Tumor-associated antigens suitable for use in the subject invention include both 35 mutated and non-mutated molecules which may be indicative of single tumor type, shared among several types of tumors, and/or exclusively expressed or overexpressed in tumor cells in comparison with normal cells. In addition to proteins and glycoproteins, tumor-specific patterns of expression of carbohydrates, gangliosides, glycolipids and mucins have also been documented. Moingeon, *supra*. Exemplary tumor-associated antigens for use in the subject

cancer vaccines include protein products of oncogenes, tumor suppressor genes and other genes with mutations or rearrangements unique to tumor cells, reactivated embryonic gene products, oncofetal antigens, tissue-specific (but not tumor-specific) differentiation antigens, growth factor receptors, cell surface carbohydrate residues, foreign viral proteins and a number 5 of other self proteins.

[00134] Specific embodiments of tumor-associated antigens include, e.g., mutated antigens such as the protein products of the Ras p21 protooncogenes, tumor suppressor p53 and HER-2/neu and BCR-abl oncogenes, as well as CDK4, MUM1, Caspase 8, and Beta catenin; 10 overexpressed antigens such as galectin 4, galectin 9, carbonic anhydrase, Aldolase A, PRAME, Her2/neu, ErbB-2 and KSA, oncofetal antigens such as alpha fetoprotein (AFP), human chorionic gonadotropin (hCG); self antigens such as carcinoembryonic antigen (CEA) and melanocyte differentiation antigens such as Mart 1/ Melan A, gp100, gp75, Tyrosinase, TRP1 and TRP2; prostate associated antigens such as PSA, PAP, PSMA, PSM-P1 and PSM- 15 P2; reactivated embryonic gene products such as MAGE 1, MAGE 3, MAGE 4, GAGE 1, GAGE 2, BAGE, RAGE, and other cancer testis antigens such as NY-ESO1, SSX2 and SCP1; mucins such as Muc-1 and Muc-2; gangliosides such as GM2, GD2 and GD3, neutral glycolipids and glycoproteins such as Lewis (y) and globo-H; and glycoproteins such as Tn, Thompson-Freidenreich antigen (TF) and sTn. Also included as tumor-associated antigens herein are 20 whole cell and tumor cell lysates as well as immunogenic portions thereof, as well as immunoglobulin idiotypes expressed on monoclonal proliferations of B lymphocytes for use against B cell lymphomas.

[00135] Tumor-associated antigens can be prepared by methods known in the art. For 25 example, these antigens can be prepared from cancer cells either by preparing crude extracts of cancer cells (e.g., as described in Cohen *et al.*, *Cancer Res.*, 54:1055 (1994)), by partially purifying the antigens, by recombinant technology, or by *de novo* synthesis of known antigens. The antigen may also be in the form of a nucleic acid encoding an antigenic peptide in a form suitable for expression in a subject and presentation to the immune system of the immunized 30 subject. Further, the antigen may be a complete antigen, or it may be a fragment of a complete antigen comprising at least one epitope.

[00136] Pathogens include, but are not limited to, infectious virus that infect mammals, and more particularly humans. Examples of infectious virus include, but are not limited to: 35 Retroviridae (e.g. human immunodeficiency viruses, such as HIV-1 (also referred to as HTLV-III, LAV or HTLV-III/LAV, or HIV-III; and other isolates, such as HIV-LP; Picornaviridae (e.g. polio viruses, hepatitis A virus; enteroviruses, human Coxsackie viruses, rhinoviruses, echoviruses); Calciviridae (e.g. strains that cause gastroenteritis); Togaviridae (e.g. equine encephalitis viruses, rubella viruses); Flaviridae (e.g. dengue viruses, encephalitis viruses,

yellow fever viruses); Coronoviridae (e.g. coronaviruses); Rhabdoviridae (e.g. vesicular stomatitis viruses, rabies viruses); Coronaviridae (e.g. coronaviruses); Rhabdoviridae (e.g. vesicular stomatitis viruses, rabies viruses); Filoviridae (e.g. ebola viruses); Paramyxoviridae (e.g. parainfluenza viruses, mumps virus, measles virus, respiratory syncytial virus);

5 Orthomyxoviridae (e.g. influenza viruses); Bungaviridae (e.g. Hantaan viruses, bunga viruses, phleboviruses and Nairo viruses); Arena viruses (hemorrhagic fever viruses); Reoviridae (e.g. reoviruses, orbiviruses and rotaviruses); Birnaviridae; Hepadnaviridae (Hepatitis B virus); Parvoviridae (parvoviruses); Papovaviridae (papilloma viruses, polyoma viruses); Adenoviridae (most adenoviruses); Herpesviridae herpes simplex virus (HSV) 1 and 2, varicella zoster virus,

10 cytomegalovirus (CMV), herpes virus; Poxviridae (variola viruses, vaccinia viruses, pox viruses); and Iridoviridae (e.g. African swine fever virus); and unclassified viruses (e.g. the etiological agents of Spongiform encephalopathies, the agent of delta hepatitis (thought to be a defective satellite of hepatitis B virus), the agents of non-A, non-B hepatitis (class 1=internally transmitted; class 2=parenterally transmitted (i.e. Hepatitis C); Norwalk and related viruses, and

15 astroviruses).

[00137] Also, gram negative and gram positive bacteria serve as antigens in vertebrate animals. Such gram positive bacteria include, but are not limited to *Pasteurella* species, *Staphylococci* species, and *Streptococcus* species. Gram negative bacteria include, but are not limited to, *Escherichia coli*, *Pseudomonas* species, and *Salmonella* species. Specific examples of infectious bacteria include but are not limited to: *Helicobacter pyloris*, *Borelia burgdorferi*, *Legionella pneumophilia*, *Mycobacteria* spp (e.g. *M. tuberculosis*, *M. avium*, *M. intracellulare*, *M. kansaii*, *M. gordonae*), *Staphylococcus aureus*, *Neisseria gonorrhoeae*, *Neisseria meningitidis*, *Listeria monocytogenes*, *Streptococcus pyogenes* (Group A

20 *Streptococcus*), *Streptococcus agalactiae* (Group B *Streptococcus*), *Streptococcus (viridans group)*, *Streptococcus faecalis*, *Streptococcus bovis*, *Streptococcus (anaerobic spp.)*, *Streptococcus pneumoniae*, pathogenic *Campylobacter* sp., *Enterococcus* sp., *Haemophilus influenzae*, *Bacillus antracis*, *corynebacterium diphtheriae*, *corynebacterium* sp., *Erysipelothrix rhusiopathiae*, *Clostridium perfringens*, *Clostridium tetani*, *Enterobacter aerogenes*, *Klebsiella*

25 *pneumoniae*, *Pasturella multocida*, *Bacteroides* sp., *Fusobacterium nucleatum*, *Streptobacillus moniliformis*, *Treponema pallidum*, *Treponema pertenue*, *Leptospira*, *Rickettsia*, and *Actinomyces israelii*.

[00138] Examples of pathogens include, but are not limited to, infectious fungi that infect

30 mammals, and more particularly humans. Examples of infectious fungi include, but are not limited to: *Cryptococcus neoformans*, *Histoplasma capsulatum*, *Coccidioides immitis*, *Blastomyces dermatitidis*, *Chlamydia trachomatis*, *Candida albicans*. Examples of infectious parasites include *Plasmodium* such as *Plasmodium falciparum*, *Plasmodium malariae*, *Plasmodium ovale*, and *Plasmodium vivax*. Other infectious organisms (i.e. protists) include

Toxoplasma gondii.

[00139] Other medically relevant microorganisms that serve as antigens in mammals and more particularly humans are described extensively in the literature, e.g., see C. G. A Thomas, 5 Medical Microbiology, Bailliere Tindall, Great Britain 1983, the entire contents of which is hereby incorporated by reference. In addition to the treatment of infectious human diseases, the compositions and methods of the present invention are useful for treating infections of nonhuman mammals. Specific examples of pathogens and antigens

10 [00140] In preferred embodiments, "treatment", "treat", "treating" as used herein with reference to infectious pathogens, refers to a prophylactic treatment which increases the resistance of a subject to infection with a pathogen or decreases the likelihood that the subject will become infected with the pathogen; and/or treatment after the subject has become infected in order to fight the infection, e.g., reduce or eliminate the infection or prevent it from becoming worse.

15 Many vaccines for the treatment of non-human mammals are disclosed in Bennett, K. Compendium of Veterinary Products, 3rd ed. North American Compendiums, Inc., 1995. As discussed above, antigens include infectious microbes such as virus, bacteria, parasites and fungi and fragments thereof, derived from natural sources or synthetically. Infectious virus of both human and non-human mammals, include retroviruses, RNA viruses, and DNA viruses.

20 This group of retroviruses includes both simple retroviruses and complex retroviruses. The simple retroviruses include the subgroups of B-type retroviruses, C-type retroviruses and D-type retroviruses. An example of a B-type retrovirus is mouse mammary tumor virus ("MMTV"). The C-type retroviruses include subgroups C-type group A (including Rous sarcoma virus ("RSV"), avian leukemia virus ("ALV"), and avian myeloblastosis virus ("AMV")) and C-type 25 group B (including murine leukemia virus ("MLV"), feline leukemia virus ("FeLV"), murine sarcoma virus ("MSV"), gibbon ape leukemia virus ("GALV"), spleen necrosis virus ("SNV"), reticuloendotheliosis virus ("RV") and simian sarcoma virus ("SSV"). The D-type retroviruses include Mason-Pfizer monkey virus ("MPMV") and simian retrovirus type 1 ("SRV-1"). The complex retroviruses include the subgroups of lentiviruses, T-cell leukemia viruses and the 30 foamy viruses. Lentiviruses include HIV-1, but also include HIV-2, SIV, Visna virus, feline immunodeficiency virus ("FIV"), and equine infectious anemia virus ("EIAV"). The T-cell leukemia viruses include HTLV-1, HTLV-II, simian T-cell leukemia virus ("STLV"), and bovine leukemia virus ("BLV"). The foamy viruses include human foamy virus ("HFV"), simian foamy virus ("SFV") and bovine foamy virus ("BFV").

35 [00141] Polypeptides of bacterial pathogens include but are not limited to an iron-regulated outer membrane protein, ("IOMP"), an outer membrane protein ("OMP"), and an A-protein of *Aeromonis salmonicida* which causes furunculosis, p57 protein of *Renibacterium salmoninarum* which causes bacterial kidney disease ("BKD"), major surface associated antigen ("msa"), a

surface expressed cytotoxin ("mpr"), a surface expressed hemolysin ("ish"), and a flagellar antigen of Yersiniosis; an extracellular protein ("ECP"), an iron-regulated outer membrane protein ("IOMP"), and a structural protein of Pasteurellosis; an OMP and a flagellar protein of *Vibrios anguillarum* and *V. ordalii*; a flagellar protein, an OMP protein, aroA, and purA of *5 Edwardsiella ictaluri* and *E. tarda*; and surface antigen of *Ichthyophthirius*; and a structural and regulatory protein of *Cytophaga columnari*; and a structural and regulatory protein of *Rickettsia*.

[00142] Polypeptides of a parasitic pathogen include but are not limited to the surface antigens
10 of *Ichthyophthirius*.

An "allergen" refers to a substance (antigen) that can induce an allergic or asthmatic response in a susceptible subject. The list of allergens is enormous and can include pollens, insect venoms, animal dander dust, fungal spores and drugs (e.g. penicillin). Examples of natural,
15 animal and plant allergens include but are not limited to proteins specific to the following genuses: Canine (*Canis familiaris*); Dermatophagoïdes (e.g. *Dermatophagoïdes farinae*); *Felis* (*Felis domesticus*); Ambrosia (*Ambrosia artemisiifolia*); *Lolium* (e.g. *Lolium perenne* or *Lolium multiflorum*); *Cryptomeria* (*Cryptomeria japonica*); *Alternaria* (*Alternaria alternata*); Alder; *Alnus* (*Alnus glutinosa*); *Betula* (*Betula verrucosa*); *Quercus* (*Quercus alba*); *Olea* (*Olea europaea*);
20 *Artemisia* (*Artemisia vulgaris*); *Plantago* (e.g. *Plantago lanceolata*); *Parietaria* (e.g. *Parietaria officinalis* or *Parietaria judaica*); *Blattella* (e.g. *Blattella germanica*); *Apis* (e.g. *Apis mellifera*); *Cupressus* (e.g. *Cupressus sempervirens*, *Cupressus arizonica* and *Cupressus macrocarpa*); *Juniperus* (e.g. *Juniperus sabina*, *Juniperus virginiana*, *Juniperus communis* and *Juniperus ashei*); *Thuya* (e.g. *Thuya orientalis*); *Chamaecyparis* (e.g. *Chamaecyparis obtusa*); *Periplaneta*
25 (e.g. *Periplaneta americana*); *Agropyron* (e.g. *Agropyron repens*); *Secale* (e.g. *Secale cereale*); *Triticum* (e.g. *Triticum aestivum*); *Dactylis* (e.g. *Dactylis glomerata*); *Festuca* (e.g. *Festuca elatior*); *Poa* (e.g. *Poa pratensis* or *Poa compressa*); *Avena* (e.g. *Avena sativa*); *Holcus* (e.g. *Holcus lanatus*); *Anthoxanthum* (e.g. *Anthoxanthum odoratum*); *Arrhenatherum* (e.g. *Arrhenatherum elatius*); *Agrostis* (e.g. *Agrostis alba*); *Phleum* (e.g. *Phleum pratense*); *Phalaris*
30 (e.g. *Phalaris arundinacea*); *Paspalum* (e.g. *Paspalum notatum*); *Sorghum* (e.g. *Sorghum halepense*); and *Bromus* (e.g. *Bromus inermis*).

[00143] Other Drug Components

Some preferred embodiments of the invention further comprise other therapeutic agents, e.g.,
35 drugs or bioactive agents. These additional components may provide direct additional therapeutic benefit or additional immune-stimulating benefits. A wide variety of therapeutic compounds may be delivered by the compositions and methods of the present invention. Examples of therapeutic compounds include, but are not limited to, nucleic acids, proteins, peptides, oncolytics, anti-infectives, anxiolytics, psychotropics, immunomodulators, ionotropes,

toxins such as gelonin and inhibitors of eucaryotic protein synthesis, and the like. Preferred therapeutic compounds for entrapment in the liposomes of the present invention are those which are lipophilic cations. Among these are therapeutic agents of the class of lipophilic molecules which are able to partition into a lipid bilayer phase of a liposome, and which

5 therefore are able to associate with the liposomes in a membrane form. Further examples of therapeutic compounds include, but are not limited to, prostaglandins, amphotericin B, methotrexate, cisplatin and derivatives, progesterone, testosterone, estradiol, doxorubicin, epirubicin, beclomethasone and esters, vitamin E, cortisone, dexamethasone and esters, betamethasone valerate and other steroids, the fluorinated quinolone antibacterial ciprofloxacin

10 and its derivatives, and alkaloid compounds and their derivatives. Among the alkaloid derivatives are swainsonine and members of the vinca alkaloids and their semisynthetic derivatives, such as, e.g., vinblastine, vincristine, vindesine, etoposide, etoposide phosphate, and teniposide. Among this group, vinblastine and vincristine, and swainsonine are particularly preferred. Swainsonine (Creaven and Mihich, *Semin. Oncol.* 4:147 (1977) has the capacity to

15 stimulate bone marrow proliferation (White and Olden, *Cancer Commun.* 3:83 (1991)). Swainsonine also stimulates the production of multiple cytokines including IL-1, IL-2, TNF, GM-CSF and interferons (Newton, *Cancer Commun.* 1:373 (1989); Olden, K., *J. Natl. Cancer Inst.*, 83:1149 (1991)). Further Swainsonine reportedly induces B- and T-cell immunity, natural killer T-cell and macrophage-induced destruction of tumor cells *in vitro*, and when combined with

20 interferon, has direct anti-tumor activity against colon cancer and melanoma cancers *in vivo* (Dennis, J., *Cancer Res.*, 50:1867 (1990); Olden, K., *Pharm. Ther.* 44:85 (1989); White and Olden, *Anticancer Res.*, 10:1515 (1990)). Other alkaloids useful in the compositions and methods of the present invention include, but are not limited to, paclitaxel (taxol) and synthetic derivatives thereof. Additional drug components, include but are not limited to, any bioactive

25 agents known in the art which can be incorporated into lipid particles.

[00144] These additional drug components may be encapsulated or otherwise associated the lipid particles described herein. Alternatively, the compositions of the invention may include drugs or bioactive agents that are not associated with the lipid-nucleic acid particle. Such drugs

30 or bioactive agents may be in separate lipid carriers or co-administered.

Manufacturing of Compositions

[00145] Manufacturing the compositions of the invention may be accomplished by any technique, but most preferred are the ethanol dialysis or detergent dialysis methods detailed in

35 the following publications, patents, and applications each incorporated herein by reference: U.S. Pat. Ser. No. 5,705,385; U.S. Pat. No. 5,976,567; U.S. Pat. Appln. No. 09/140,476; U.S. Pat. No. 5,981,501; U.S. Pat. No. 6,287,591; Int. Publ. No. WO 96/40964; and Int. Publ. No. WO 98/51278. These manufacturing methods provide for small and large scale manufacturing of immunostimulatory compositions comprising therapeutic agents encapsulated in a lipid

particle, preferably lipid-nucleic acid particles. The methods also generate such particles with excellent pharmaceutical characteristics.

[00146] Vaccine compositions of the present invention may be prepared by adding a target antigen (to which the immune response is desired). Means of incorporating antigens are well known in the art and include, for example: 1) passive encapsulation of the antigen during the formulation process (e.g., the antigen can be added to the solution containing the ODN); 2) addition of glycolipids and other antigenic lipids to an ethanol lipid mixture and formulated using the ethanol-based protocols described herein; 3) insertion into the lipid vesicle (e.g., antigen-lipid can be added into formed lipid vesicles by incubating the vesicles with antigen-lipid micelles); and 4) the antigen can be added post-formulation (e.g., coupling in which a lipid with a linker moiety is included into formulated particle, and the linker is activated post formulation to couple a desired antigen). Standard coupling and cross-linking methodologies are well known in the art. An alternative preparation incorporates the antigen into a lipid-particle which does not contain a nucleic acid, and these particles are mixed with lipid-nucleic acid particles prior to administration to the subject.

Characterization of Compositions Used in the Methods of the Present Invention

[00147] Preferred characteristics of the compositions used in the methods of the present invention are as follows.

[00148] The preferred lipid-nucleic acid particles of the invention comprise a lipid membrane (generally a phospholipid bilayer) exterior which fully encapsulates an interior space. These particles, also sometimes herein called lipid membrane vesicles, are small particles with mean diameter 50-200 nm, preferably 60-130 nm. Most preferred for intravenous administrations are particles of a relatively uniform size wherein 95% of particles are within 30 nm of the mean. The nucleic acid and other bioactive agents are contained in the interior space, or associated with an interior surface of the encapsulating membrane.

[00149] "Fully encapsulated" as used herein indicates that the nucleic acid in the particles is not significantly degraded after exposure to serum or a nuclease assay that would significantly degrade free DNA. In a fully encapsulated system, preferably less than 25% of particle nucleic acid is degraded in a treatment that would normally degrade 100% of free nucleic acid, more preferably less than 10% and most preferably less than 5% of the particle nucleic acid is degraded. Alternatively, full encapsulation may be determined by an Oligreen™ assay. Fully encapsulated also suggests that the particles are serum stable, that is, that they do not rapidly decompose into their component parts upon *in vivo* administration.

[00150] These characteristics of the compositions of the present invention distinguish the preferred particles of the invention from lipid-nucleic acid aggregates (also known as cationic complexes or lipoplexes) such as DOTMA/DOPE (LIPOFECTIN™) formulations. These aggregates are generally much larger (>250 nm) diameter, they do not competently withstand 5 nuclease digestion. They generally decompose upon *in vivo* administration. Lipid-nucleic acid formulations comprising cationic lipid-nucleic acid aggregates with weak antigens, as described above, may provide suitable vaccines for local and regional applications, such as intra-muscular, intra-peritoneal and intrathecal administrations, and more preferably intranasal administration.

10 [00151] The liposomal particles of the invention can be formulated at a wide range of drug:lipid ratios. "Drug to lipid ratio" as used herein refers to the amount of therapeutic nucleic acid (*i.e.*, the amount of nucleic acid which is encapsulated and which will not be rapidly degraded upon exposure to the blood) in a defined volume of preparation divided by the amount of lipid in the 15 same volume. This may be determined on a mole per mole basis or on a weight per weight basis, or on a weight per mole basis. Drug to lipid ratio may determine the lipid dose that is associated with a given dose of nucleic acid. In a preferred embodiment, the compositions of the present invention have a drug:lipid ratio in the range of about 0.01 to 0.25 (wt/wt).

20 **Uses of the Compositions and Methods of the Present Invention**

[00152] The combination of B lymphocytes and T lymphocytes establish the underlying operation of the humoral and cellular immune responses, respectively. The humoral and cellular immune responses each proceed by activation of their respective cell types in response to stimulation from an antigen and the consequent secretions of various cytokines. The 25 presentation of antigenic peptide to naïve CD4+ T helper cells causes the cells to differentiate into two distinct subsets of helper cells (Th-1 and Th-2) which can be distinguished by their function and cytokine expression profiles. Mosman *et al.*, *Annu. Rev. Immunol.*, 7:145-173 (1989); Paul *et al.*, *Cell*, 76: 241-251 (1994); O'Garra, *Immunity*, 8:275-283 (1998).

30 [00153] The specific patterns of cytokines secreted by the CD4+ Th cells steer the immune response to a predominantly cellular, type-1 response (including IFN- γ , IL-1, IL-2, IL-12, and TNF- α) or a mainly humoral, type-2 response (including IL-4, IL-5, IL-6, IL-9, IL-10 and IL-13). Glimcher and Murphy, *Genes Dev.*, 14:1693-1711 (2000); Abbas *et al.*, *Nature*, 383:787-793 35 (1996). The Th-1 subset promotes both cell-mediated immunity through activation of CTL and NK cells, as well as humoral immunity characterized by immunoglobulin class switching from IgM to IgG and IgA in humans, and to IgG2a in mice. Th-1 responses may also be associated with delayed-type hypersensitivity and autoimmune disease. The Th-2 subset induces primarily humoral immunity and induces class switching to IgG1 and IgE in humans. The antibody isotypes associated with Th-1 responses generally have good neutralizing and opsonizing

capabilities whereas those associated with Th-2 responses are generally more associated with allergic responses.

[00154] As demonstrated herein, the subject immunostimulatory compositions are capable of stimulating a strong, Th-1 biased immune response to antigenic stimulation, e.g., microbial antigens and tumor-associated antigens, and can enhance both the cellular and the humoral components of the host immune response. Thus, the immunostimulatory compositions described herein find use as adjuvants in methods of inducing Th-1 biased immune responses in general, and in vaccines directed to specific antigen(s) of interest. Also provided herein are 10 methods for improving the maturation of the humoral response as well as methods for increasing antibody isotype switching in response to antigenic stimulation.

[00155] These immune responses can be measured in many ways including but not limited to activation, proliferation or differentiation of cells of the immune system (e.g., B cells, T cells, 15 APCs, such as dendritic cells or macrophages, NK cells, NKT cells etc.); up-regulated or down-regulated expression of markers; cytokine secretion; stimulation of or increase in IgA, IgM, or IgG titer; isotype class switching, and splenomegaly (including increased spleen cellularity). The presence of a Th-1 biased immune response in particular can be determined directly by the induction of Th-1 cytokines (e.g., IFN- γ , IL-12) and antigen-specific CD8+ CTL. Thus, if Th-1 20 cytokines or CTL are induced, Th-1 biased immune responses are induced according to the invention. Similarly, enhanced humoral responses and improvements in the maturation of the humoral response are indicated by detecting the isotype of type-1 antigen-specific antibodies that are induced (e.g., IgG2a, IgG1 in mice, IgG and IgA in humans), and determining if isotype switching has occurred, e.g., IgM to IgG or IgA, as exemplified herein. If increased isotype 25 switching has occurred in comparison with alternative adjuvants, enhanced humoral immune responses are induced according to the invention.

[00156] In a preferred embodiment, the methods of the present invention comprise stimulating a Th1-baised immune response to antigenic stimulation by administering to the subject an 30 effective amount of an immunostimulatory composition comprising an LNA formulation including a methylated nucleic acid. Vaccines are also provided comprising such LNA formulations in combination with one or more epitopes of one or more antigens of interest. Preferably the antigen is associated with the LNA particle, and most preferably a plurality of 35 antigens are employed. In one embodiment, the ODN comprises a PS or other modified, non-phosphodiester backbone. Alternative adjuvants that induce Th1 responses include but are not limited to MPL, MDP, ISCOMS, IL-12, IFN- γ , and SB-AS2.

[00157] In a further embodiment, the compositions and methods of the present invention can be used to modulate the level of a cytokine. "Modulate" as used herein with reference to a

cytokine may refer to the suppression of expression of a particular cytokine when lower levels are desired, or augmentation of the expression of a particular cytokine when higher levels are desired. Modulation of a particular cytokine can occur locally or systemically. In a preferred embodiment, the compositions and methods of the present invention can be used to activate 5 macrophages and dendritic cells to secrete cytokines. In general, Th1-type cytokines can be induced, and thus the immunostimulatory compositions of the present invention can promote a Th1 type antigen-specific immune response including cytotoxic T-cells.

Indications, Administration and Dosages

10 [00158] The compositions and methods of the present invention are indicated for use in any patient or organism having a need for immune system stimulation. Such a need encompasses, but is not limited to, most medical fields, such as oncology, inflammation, arthritis & rheumatology, immuno-deficiency disorders. One skilled in the art can select appropriate indications to test for efficacy based on the disclosure herein. In a preferred embodiment, the 15 compositions and methods of the invention are used to treat a neoplasia (any neoplastic cell growth which is pathological or potentially pathological) such as the neoplasia described in the Examples below.

20 [00159] Administration of the compositions of the invention to a subject may be by any method including *in vivo* or *ex vivo* methods. *In vivo* methods can include local, regional or systemic applications. In a preferred embodiment, the compositions are administered intravenously such that particles are accessible to B cells, macrophages or splenocytes in a patient, and/or the particle can stimulate lymphocyte proliferation, resulting in secretion of IL-6, IL-12, IFNg and/or IgM in said patient.

25 [00160] Vaccine compositions of the present invention may be administered by any known route of administration. In one embodiment, the compositions of the present invention are administered via intravenous injection. In another embodiment, intramuscular or subcutaneous injection is employed and in this manner larger-sized (150-300 nm) lipid particles can be used.

30 Consequently, the need for costly extrusion steps can be reduced or eliminated, and since the particles do not need to circulate, the selection of lipid components can be biased in favor of less expensive materials. For example, the amount of Chol can be reduced, DSPC can be replaced with something less rigid (e.g., POPC or DMPC), and PEG-lipids can be replaced with less expensive PEG-acyl chains. In a still further embodiment, the compositions of the present 35 invention are administered via the respiratory tract, e.g., by intratracheal instillation or intranasal inhalation.

[00161] One skilled in the art would know how to identify possible toxicities of formulations, for example, complement activation, coagulation, renal toxicities, liver enzyme assays, etc. Such

toxicities may differ between organisms.

[00162] Pharmaceutical preparations of compositions usually employ additional carriers to improve or assist the delivery modality. Typically, compositions of the invention will be 5 administered in a physiologically-acceptable carrier such as normal saline or phosphate buffer selected in accordance with standard pharmaceutical practice. Other suitable carriers include water, 0.9% saline, 0.3% glycine, and the like, including glycoproteins for enhanced stability, such as albumin, lipoprotein, globulin, etc.

10 [00163] Dosages of lipid-nucleic acid formulations depend on the desired lipid dosage, the desired nucleic acid dosage, and the drug:lipid ratio of the composition. One skilled in the art can select proper dosages based on the information provided herein.

[00164] Immunotherapy or vaccination protocols for priming, boosting, and maintenance of 15 immunity are well known in the art and further described below. In particular, one skilled in the art would know how to calculate dosage amounts for a subject, particularly a mammal, and more particularly a human, based on the dosage amounts described herein. Specific conversion factors for converting dosage amounts from one animal to another (e.g., from mouse to human) are well known in the art and are fully described, e.g., on the Food and Drug 20 Administration Web site at: www.fda.gov/cder/cancer/animalframe.htm (in the oncology tools section), incorporated herein by reference. As compared to known immunostimulatory compositions having free nucleic acids, the immunostimulatory compositions and methods of the present invention may utilize reduced amounts of nucleic acids to stimulate enhanced immune responses *in vivo*.

25 [00165] The amount of nucleic acids in the formulations of the present invention will generally vary between about 0.001-60 mg/kg (mg nucleic acids per kg body weight of a mouse per dose). In preferred embodiments for intravenous (i.v.) administration, the compositions and methods of the present invention utilize about 1-50 mg/kg, more preferably about 5-20 mg/kg.

30 In preferred embodiments for subcutaneous (s.c.) administration, the compositions and methods of the present invention utilize about 1-10 mg/kg, and more preferably about 1-5 mg/kg, usually about about 3-5 mg/kg. The amount of antigen associated with the lipid particles of the present invention is preferably about 0.04-40 mg/kg, and more preferably about 0.04-4 mg/kg. As described above, one skilled in the art could readily determine suitable 35 dosage amounts for other mammals given the dosage amounts described herein, based on the well-known conversion factors identified above and further empirical testing.

[00166] The formulations of the invention may be administered in pharmaceutically acceptable solutions, which may routinely contain pharmaceutically acceptable concentrations of salt,

buffering agents, preservatives, compatible carriers, adjuvants, and optionally other therapeutic ingredients.

[00167] For use in therapy, an effective amount of the immunostimulatory compositions of the present invention can be administered to a subject by any mode allowing uptake by the appropriate target cells. "Administering" the immunostimulatory composition of the present invention may be accomplished by any means known to the skilled artisan. Preferred routes of administration include but are not limited to parenteral injection (subcutaneous, intradermal, intravenous, parenteral, intraperitoneal, intrathecal, etc.), as well as mucosal, intranasal, intratracheal, inhalation, and intrarectal, intravaginal; or oral, transdermal (e.g., via a patch). An injection may be in a bolus or a continuous infusion.

[00168] For example, the immunostimulatory compositions of the present invention can be administered by intramuscular or intradermal injection, or other parenteral means, or by biostatic "gene-gun" application to the epidermis. The immunostimulatory compositions of the present invention may also be administered, for example, by inhalation, topically, intravenously, orally, implantation, rectally, or vaginally. Suitable liquid or solid pharmaceutical preparation forms are, for example, aqueous or saline solutions for injection or inhalation, encochleated, coated onto microscopic gold particles, and nebulized. For a brief review of present methods for drug delivery, see Langer, *Science* 249:1527-1533, 1990, which is incorporated herein by reference.

[00169] The pharmaceutical compositions are preferably prepared and administered in dose units. Liquid dose units are vials or ampoules for injection or other parenteral administration. Solid dose units are tablets, capsules and suppositories. For treatment of a patient, depending on activity of the compound, manner of administration, purpose of the immunization (i.e., prophylactic or therapeutic), nature and severity of the disorder, age and body weight of the patient, different doses may be necessary. The administration of a given dose can be carried out both by single administration in the form of an individual dose unit or else several smaller dose units. Multiple administration of doses at specific intervals of weeks or months apart is usual for boosting the antigen-specific responses.

[00170] Suitable buffering agents include: acetic acid and a salt (1-2% w/v); citric acid and a salt (1-3% w/v); boric acid and a salt (0.5-2.5% w/v); and phosphoric acid and a salt (0.8-2% w/v). Suitable preservatives include benzalkonium chloride (0.003-0.03% w/v); chlorobutanol (0.3-0.9% w/v); parabens (0.01-0.25% w/v) and thimerosal (0.004-0.02% w/v).

[00171] In preferred embodiments, the immunostimulatory compositions of the present invention contain an effective amount of a combination of adjuvants and antigens optionally included in a pharmaceutically-acceptable carrier. "Pharmaceutically-acceptable carrier" as

used herein refers to one or more compatible solid or liquid filler, dilutants or encapsulating substances which are suitable for administration to a human or other mammal. "Carrier" as used herein refers to an organic or inorganic ingredient, natural or synthetic, with which the active ingredient is combined to facilitate the application. The components of the

5 immunostimulatory compositions of the present invention also are capable of being comingled with the compounds of the present invention, and with each other, in a manner such that there is no interaction which would substantially impair the desired pharmaceutical efficiency.

[00172] Compositions suitable for parenteral administration conveniently comprise sterile

10 aqueous preparations, which can be isotonic with the blood of the recipient. Among the acceptable vehicles and solvents are water, Ringer's solution, phosphate buffered saline and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose any bland fixed mineral or non-mineral oil may be employed including synthetic mono-ordi-glycerides. In addition, fatty acids such as oleic 15 acid find use in the preparation of injectables. Carrier formulations suitable for subcutaneous, intramuscular, intraperitoneal, intravenous, etc. administrations may be found in Remington's Pharmaceutical Sciences, Mack Publishing Company, Easton, Pa.

[00173] The adjuvants or antigens useful in the invention may be delivered in mixtures of more

20 than two adjuvants or antigens. A mixture may consist of several adjuvants in addition to the LNA formulations described herein.

[00174] A variety of administration routes are available. The particular mode selected will depend, of course, upon the particular adjuvants or antigen selected, the age and general

25 health status of the subject, the particular condition being treated and the dosage required for therapeutic efficacy. The methods of this invention, generally speaking, may be practiced using any mode of administration that is medically acceptable, meaning any mode that produces effective levels of an immune response without causing clinically unacceptable adverse effects. Preferred modes of administration are discussed above.

30

[00175] The compositions may conveniently be presented in unit dosage form and may be prepared by any of the methods well known in the art of pharmacy. All methods include the step of bringing the compounds into association with a carrier which constitutes one or more accessory ingredients. In general, the compositions are prepared by uniformly and intimately 35 bringing the compounds into association with a liquid carrier, a finely divided solid carrier, or both, and then, if necessary, shaping the product.

[00176] Other delivery systems can include time-release, delayed release or sustained release delivery systems. Such systems can avoid repeated administrations of the compounds,

increasing convenience to the subject and the physician. Many types of release delivery systems are available and known to those of ordinary skill in the art. They include polymer base systems such as poly(lactide-glycolide), copolyoxalates, polycaprolactones, polyesteramides, polyorthoesters, polyhydroxybutyric acid, and polyanhydrides. Microcapsules 5 of the foregoing polymers containing drugs are described in, for example, U.S. Pat. No. 5,075,109. Delivery systems also include non-polymer systems that are: lipids including sterols such as cholesterol, cholesterol esters and fatty acids or neutral fats such as mono-di-and tri-glycerides; hydrogel release systems; sylastic systems; peptide based systems; wax coatings; compressed tablets using conventional binders and excipients; partially fused implants; and the 10 like. Specific examples include, but are not limited to: (a) erosional systems in which an agent of the invention is contained in a form within a matrix such as those described in U.S. Pat. Nos. 4,452,775, 4,675,189, and 5,736,152, and (b) diffusional systems in which an active component permeates at a controlled rate from a polymer such as described in U.S. Pat. Nos. 3,854,480, 5,133,974 and 5,407,686. In addition, pump-based hardware delivery systems can be used, 15 some of which are adapted for implantation.

[00177] Turning now to certain particular aspects of the present invention, one aspect arises from the recognition that *in vitro* data showing effects of oligonucleotides on stimulation of various leukocyte markers, including activation markers, are not determinative of whether a free 20 oligonucleotide is, in-fact, immunostimulatory *in vivo* or will have therapeutic efficacy.

EXPERIMENTAL

Experimental Details

25 [00178] Mice. Female, Balb/c or C57/BL6 ("B6") mice (6-8 weeks) were purchased from Harlan-Sprague Dawley (Indianapolis, IN). All animals were quarantined for one week prior to use. All studies were conducted in accordance with the guidelines established by the Canadian Council on Animal Care (CCAC) and the Institutional Animal Care and User Committee (IACUC).

30 [00179] Peptides. Peptides were obtained from Commonwealth Biotechnologies and were >95% pure as determined by HPLC. QC analyses were obtained with each peptide.

[00180] Oligonucleotide formulations. Oligonucleotides were obtained from Avecia or Proligo 35 or Trilink. All ODN were rehydrated in sterile water and diluted to the appropriate concentrations in sterile DPBS, pH 7.2.

[00181] LNA formulations. LNA formulations were prepared as follows. Initial ODN:lipid ratios were 0.25, w/w. For phosphodiester formulations, 20 mM citrate buffer was used in place of

300 mM citrate buffer to dissolve the ODN. Failure to do this results in considerably diminished encapsulation efficiency (i.e. ~ 3% final).

[00182] Vaccinations. B6 mice were vaccinated subcutaneously (SC) with 100 ml per injection.

5. Dosing schedules were either q7dx2 or q4dx4 and are indicated in the individual figure legends. CFA mixtures were very viscous and had to be given with a larger gauge syringe. For humoral studies, a prime (day 0) and boost (day 14) strategy was used and blood samples were collected at various times by tail nicking.

10 [00183] Efficacy studies. E.G7-OVA or EL-4 thymoma cell lines were used throughout the study. These cells were cultured *in vitro* according to established methods. For tumor studies, 2.5 x 10⁶ E.G7-OVA cells were injected subcutaneously in 50 ml of PBS containing 1% FCS. Tumor measurements were made by repeated perpendicular measurements of tumor dimensions and using the formula:

15

$$\text{Volume (mm}^3\text{)} = (\text{L} \times \text{W} \times \text{H})\text{p/6}$$

[00184] TRP-2 and gp100 studies were conducted using the B16/BL6 murine melanoma model. B16 cells (1.0x10⁵) were injected IV in a volume of 200 ml. Typically, animals were

20 vaccinated weekly for 2-3 injections and B16 cells were then administered 1-2 days after the final vaccination. Animals were terminated between days 14-18 post-B16 injection, lungs were removed, and metastases were counted using a stereomicroscope.

[00185] Flow cytometry analysis. Antigen specific T cells were determined in vaccinated mice

25 at various times using either MHC tetramers (Beckman Coulter) or the Dimer X reagent (Pharmingen). Antibodies against CD8a, B220, and CD4 were used to identify cell populations of interest and gate out unwanted populations. Single cell suspensions of spleen and lymph nodes were prepared according to protocols outlined in Current Protocols in Immunology. Antibody stains were done on 1 x 10⁶ cells in 96 well plates kept at 4°C. Generally, 0.2 mg of

30 each antibody was used per 1 x 10⁶ cells.

Example 1

In vitro vs. In vivo Activation of Leukocytes in Whole Blood Cells by Exposure to Free or Encapsulated Oligonucleotides

35

[00186] In order to demonstrate the effectiveness of an *in vitro* assay for predicting immune stimulation *in vivo* a comparison of CD69 expression is shown in Figures 1 and 2. CD69 is a cell activation marker, which quantifies the activation of NK cells, B cells and monocytes. Expression of CD69 on NK cells indicates cell activation and production of IFN- γ , which is

important to inducing a Th-1 immune response. Free and encapsulated ODN1 and 2 were tested in vitro and in vivo for their ability to induce CD69 expression. A dose of 0.1 mg/ml of ODN2 and 10mg/ml ODN1 were used in vitro and 10mg/kg of ODN2 and 20mg/kg of ODN1 in vivo. Each oligonucleotide was encapsulated in a lipid particle composed of

5 POPC:CHOL:DODMA:PEGDMG in a ratio of 20:45:25:10.

[00187] Figure 1 illustrates the in vitro stimulation of leukocytes bearing the activation marker CD69 from treating mouse whole blood with free oligonucleotides and encapsulated oligonucleotides, specifically ODN1 and 2. When mouse whole blood was treated in vitro with

10 free oligonucleotides there was a dose responsive increase in the amount of CD69 positive B-cells, monocytes and to some extent, NK cells, according to the amount of free oligonucleotide used 15 hours after treatment. In this in vitro assay, free ODN2 caused much greater stimulation of CD69 than free ODN1. However, when these same oligonucleotides were encapsulated in a lipid vesicle, the in vitro stimulation of CD69 production on these same cell types was reduced or abolished altogether.

15 [00188] When the same oligonucleotides were tested in vivo, however, surprising results were obtained. Figure 2 illustrates that in vivo treatment of ICR mice by injection with encapsulated or free oligonucleotides produces results that are contrary to those obtained in vitro. This figure 20 clearly demonstrates that in vivo, the lipid encapsulated oligonucleotides were more effective than the free oligonucleotides in stimulating the CD69 marker on the same cell types at 16 and 24 hours after injection. The results show that in vitro data is not sufficient for determining whether an oligonucleotide will be immunostimulatory in vivo. Moreover, Figures 1 and 2 suggest that lipid encapsulation is an important factor in determining whether an oligonucleotide 25 would be effective in vivo. The in vivo results show that encapsulated ODN1 and ODN2 were both able to stimulate production of CD69 on NK cells, whereas the in vitro results indicated lipid encapsulation of ODN1 and 2 actually reduced the stimulation of CD69 on NK cells below the control level.

30 [00189] The foregoing in vivo results show that free oligonucleotides are not necessarily immunostimulatory unless they are encapsulated in a lipid vesicle as measured by stimulation of CD69 bearing cells in vivo. This is true even though stimulation of CD69 was observed in vitro by free ODN. Further, the results indicate that an encapsulated oligonucleotide may be effective in vivo even though it is not shown to be effective in vitro.

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Example 2

In vivo Dendritic Cell Activation with Methylated Oligonucleotides

[00190] As discussed in the Background section above, the prior art teaches that methylated CpG oligonucleotides are generally not effective, or less effective in comparison to unmethylated CpG oligonucleotides in stimulating immune responses whether measured in vitro or in vivo. United States Patent No. 6,429,199 discloses that methylated oligonucleotides 5 did not enhance the expression of CD40 on NK cells or human B cells, nor did they show any improved survival of dendritic cells, which are the major antigen presenting cells involved in humoral and cellular immunity in a Th-1 response. Further, the methylated CpG oligonucleotides disclosed were inactive in improving survival, differentiation, activation or 10 maturation of dendritic cells in vitro. Similarly, the in vitro PBMC results disclosed in WO 02/069369 did not demonstrate any activity of methylated oligonucleotides on dendritic cells.

[00191] In contrast, the present invention shows that methylated oligonucleotides are at least as effective and typically more effective at inducing proliferation of dendritic cells than unmethylated oligonucleotides. The counterpart of unmethylated ODN1 was made where the 15 cytosine residue of the single CpG dinucleotide sequence was methylated and is referred to as ODN1m.

[00192] In order to demonstrate that methylated oligonucleotides are capable of stimulating dendritic cells each of ODN1 and ODN1m were encapsulated in a lipid vesicle 20 comprising POPC:CHOL:DODMA:PEGDMG in a ratio of 20:45:25:10. PBS was used as a control. The results of this experiment are shown in Figure 3.

[00193] Figure 3 clearly illustrates the ability of encapsulated ODN1m to activate dendritic cells. Furthermore, when encapsulated in a lipid vesicle, methylated ODN1m was more active than its 25 unmethylated counterpart ODN1 in stimulating activation of dendritic cells in vivo. Dendritic cell activation was measured by the percentage of IFN- γ secreting cells. These cells were labeled with an antibody to indicate the cell type and only those having the dendritic cell marker were included in this measurement.

30 [00194] In order to demonstrate the expansion of dendritic cells resulting from the administration of lipid-encapsulated unmethylated and methylated CpG oligonucleotides, cells from the blood, spleen and lymph nodes were analyzed for activation and expansion of dendritic cell populations. ICR mice were immunized with a single intravenous injection of encapsulated oligonucleotides at a dose of 20 mg/kg and the control ICR mouse was injected with PBS. 35 Cells for each of the spleen, blood and lymph nodes were isolated at various time points, as shown in Figures 4 and 5, and the amount of dendritic cell expansion and activation was measured through the use of dendritic cell markers CD11c and DEC205. Each of these markers are specific to dendritic cells though they may represent different cell sub-populations. In each of Figures 4 and 5 the control was plotted as equivalent to 100% and the effect of

ODN1m for each backbone configuration was plotted as a percentage of that control. Figure 4A illustrates in each of the panels that the methylated ODN1m stimulated the expansion of CD11c positive dendritic cells in spleen cells and whole blood cells but not in lymphoid tissue as measured against the control. Both the PO and PS backbones for ODN1m showed dendritic cell expansion. Figure 4B shows in each of the panels that methylated ODN1m also stimulated the expansion of DEC205 positive dendritic cells in spleen cells and lymphoid tissue but not in whole blood cells as measured against the control.

[00195] Figure 5 also demonstrates the activation of dendritic cells. On collection of the 10 samples the cells were first analyzed by flow cytometry for the co-expression of CD86, which indicates cell activation, and the dendritic cell phenotype markers CD11c and DEC205. The percentage of activated dendritic cells was plotted against a PBS control equivalent to 100%. Figure 5, each of the panels show that the methylated ODN1m induced CD86 expression on 15 CD11c positive dendritic cells when the oligonucleotide was lipid encapsulated. Similar results are shown when measuring DEC205 positive dendritic cells in Figure 5 panels.

[00196] The data in Figures 4 and 5 therefore refutes the statements in United States Patent 20 No. 6,429,199 which teaches that methylated CpG oligonucleotides are inactive in improving survival, differentiation, activation or maturation of dendritic cells.

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Example 3

The Effect of PS and PO Backbone Configurations on Plasma Cytokine Levels

[00197] As noted above, with non-lipid encapsulated oligonucleotides the backbone is 25 traditionally modified so as to reduce molecular degradation by nucleases. However, on encapsulation, such a modification is no longer required to prevent degradation. To establish the effect of a different phosphate backbone on cytokine stimulation induced by lipid encapsulation, mice were injected with oligonucleotides having both PO and PS backbones, and cytokine stimulation was measured over a series of points in time (as shown in Figure 6 30 and 7B) and over a sliding dosage scale as shown in Figure 7A. In this experiment ICR mice were injected i.v. with a 20mg ODN/kg dose of free PO ODN1, encapsulated PO-ODN1 and encapsulated PS-ODN1. Cytokines common to both Th-1 and Th-2 (IL-12, IL-6 and IFN- γ) and MCP-1 (a macrophage chemokine) were measured over a 24-hour time course following 35 administration. Cytokine stimulation is generally indicative of a cellular immune response as is a chemokine response in MCP-1. Oligonucleotides were encapsulated in a lipid particle composed of DSPC:Chol:DODAP:PEG-CER14 in a ratio of 20:45:25:10.

[00198] Figure 6 shows that in vivo administration of free PO-ODN1 had no effect on stimulation of IL-6, IL-12 IFN- γ or MCP-1 indicating that the oligonucleotide was likely degraded

by nucleases. It is well known in the art that a PS backbone is required when administering free oligonucleotide in order to avoid nuclease degradation. In contrast, in vivo administration of lipid encapsulated PO and PS ODN1 stimulated production of each of these cytokines and chemokine. However, Figure 6 indicates that the PO-ODN is more effective at inducing 5 cytokine and chemokine production.

[00199] Figure 7A illustrates increased IL-12 induction by treatment of ICR mice with either encapsulated PO or PS ODN14 in comparison to free ODN14 measured over a sliding dosage scale. This figure supports the conclusions drawn from Figure 6 indicating that lipid 10 encapsulation increases the effectiveness of cytokine stimulation as evidenced by an increase in IL-12 induction and that a PO-ODN is more effective at inducing a cytokine response than a PS-ODN. In fact, when administered at the same dose but in encapsulated form a 2.5 fold increase in peak plasma IL-12 is observed, along with even more dramatic increases in other cytokines such as IL-6 (1000-fold) and IFN- γ (20-fold). Figure 7A also demonstrates that a 15 lower dose of oligonucleotides is required to facilitate a cytokine response when encapsulated in comparison with free oligonucleotides administered in the absence of the lipid particles.

[00200] In order to further elucidate the difference in PO-ODN and PS-ODN Figure 7B illustrates differences in IFN- γ cytokine stimulation over time specific to PS and PO backbone 20 configurations. Treatment with encapsulated PO ODN14 stimulates a strong early induction of IFN-gamma while treatment with encapsulated PS ODN14 stimulates a smaller but still effective induction of IFN-gamma. Moreover, Figure 7B shows that over a period of days, a second large IFN- γ peak occurred when stimulating with the PS-ODN which may indicate that the immune system was primed by treatment with the encapsulated PS-ODN to respond more 25 effectively to IL-12 production, possibly through expansion of NK cells after treatment.

[00201] Figure 22 similarly demonstrates the late IFN- γ peak seen in Figure 7B for the PS-ODN in comparison with the PO version of the same oligonucleotides, as discussed further in Example 4.

30 [00202] An important feature of lipid encapsulation according to the present invention is the finding that oligonucleotides having natural PO backbones can be used to stimulate an immune response whereas in the prior art, PS backbones are required for effective in vivo activity. As shown in Example 7 below, encapsulated PO oligonucleotides may be more effective than PS 35 oligonucleotides when evaluating anti-tumor efficacy, especially where the oligonucleotide is methylated.

Example 4

Evaluation of immunostimulatory properties of CpG ODN having PS and PO backbones

[00203] When differentiating the levels of response particularly associated with ODN's having PO and PS backbones, a further aspect is the analysis of the type of response being evaluated, more specifically, whether the response is a humoral response or a cellular response. In order 5 to assess the effect of the backbones, an experiment was conducted to look at the ability of PS-ODN and PO-ODN to initiate and induce maturation (i.e. facilitate isotype switching) of a humoral immune response. The magnitude and kinetics of a humoral immune response elicited by administration of encapsulated PO-ODN and PS-ODN was compared. Each of the 10 PO-ODN and PS-ODN were administered subcutaneously at a dose of 100 mg/dose in a q14 x 2 prime-boost setting on Days 0 and 14 and assessed at 6 weeks on Day 35. The control mice were immunized at the same dose using OVA-PBS and OVA-Alum. Oligonucleotides were 15 encapsulated in a lipid particle composed of DSPC:Chol:DODMA:PEG-DMG in a ratio of 20:45:25:10.

15 [00204] It can be concluded that although both PO and PS ODNs are able to induce a humoral immune response, the nature of the response is different. Figures 8 and 9 illustrate the magnitude of the IgM and IgG response after 6 weeks for the various oligonucleotides and control in terms of absorbance. As is clearly demonstrated in the Figures, each of PS-ODN1 and PS-ODN2 produced a weak IgM response whereas PO-ODN2 produced a strong IgM 20 response. Conversely, the IgG response produced was consistently better for each of the PS-ODN tested in comparison with the same oligonucleotides having a PO backbone. This suggests that a PS-ODN produces a superior IgG response. These data indicate that while both PO and PS ODN are able to initiate a humoral immune response, the PO response does not mature as indicated by a lack of isotype switching and a preponderance of the IgM isotype. 25 On the other hand, PS-ODN are able to initiate a humoral immune response as well as induce maturation of the response as indicated by isotype switching to a dominance of IgG isotype antibodies.

[00205] This phenomenon may be related to the cytokine profiles induced by PO vs. PS ODN. 30 Figure 7(b) and Figure 22 illustrate that encapsulated PS oligonucleotides ODN1 and ODN2 produced a strong IFN- γ peak 6 days after treatment that is not produced by encapsulated PO oligonucleotides. It has been reported that cytokines such as IFN- γ result in preferential isotype switching to various IgG isotypes. Therefore, the large PS-ODN-induced late IFN- γ peak may 35 induce isotype switching from IgM to IgG isotypes while the lack of such a peak in PO-ODN-treated mice may result in no isotype switching. The basis for this reduced late IFN- γ peak with PO-ODN is not clear, but results may suggest that treatment with encapsulated PO oligonucleotides but not PS oligonucleotides causes a prior induction of type I interferons that inhibit the expression of IL-12, which is needed to promote IFN- γ expression in NK or T cells.

[00206] Similarly, Figure 6 not only shows that encapsulation of the oligonucleotides is important for stimulating the production of cytokines that lead to a Th-1 response as previously discussed, but also shows that more cytokines are produced using encapsulated PO oligonucleotides than PS oligonucleotides. This contrasts with administration of free 5 oligonucleotides as taught in the prior art, which generally shows that a PS backbone is preferred over PO oligonucleotides to prevent degradation of the oligonucleotide in vivo.

Example 5

In vivo Immunological Responses to Treatment with Oligonucleotides as measured by
10 Cytokine Induction, Tetramer Analysis and Cytotoxicity Assay (CTL)

[00207] In vivo Immunological Responses to Treatment with Oligonucleotides as measured by Cytokine Induction, Tetramer Analysis and Cytotoxicity Assay (CTL)

15 [00208] To monitor immunological response to subcutaneous immunization, antigen specific cellular immune responses were monitored using MHC Class I-tetramer analyses, cytotoxicity assays and cytokine release assays while humoral immune responses were monitored by measuring plasma antibody levels. Cellular and humoral responses were assessed in C57Bl/6 and Balb/C mice respectively (5 animals per group). For analysis of the cellular response, mice 20 were immunized subcutaneously with 3 injections on a q7d x 3 dosing regimen on Days 0, 7 and 14 at a dose of 100 mg oligonucleotide in combination with 20mg of antigen. The spleen, liver, lymph node and blood tissues were collected on Day 21. Solid tissues were mechanically dissociated and cells were processed to collect mononuclear cells. For analysis of the humoral response, animals were immunized twice on a q14d x 2 on Days 0 and 14 and blood was 25 collected on Day 35 for analysis of plasma for immunoglobulin levels. In this series of experiments oligonucleotides were encapsulated either in lipid particles composed of DSPC:Chol:DODMA:PEG-DMG or POPC:Chol:DODMA:PEG-DMG at a ratio of 20:45:25:10. All comparisons were done with like lipid particles.

30 [00209] MHC-tetramer analysis is designed to detect CD8+ve, cytotoxic T-lymphocytes that possess the appropriate T-cell receptor to allow recognition and lysis of target cells bearing the target antigen in the context of a MHC Class I complex. Isolated splenocytes from immunized animals were stained with PE-labeled MHC Class I tetramers (H2K^b) complexed with the immunodominant OVA SIINFEKL peptide as well as FITC-labeled anti-CD8 and Cy-Chrome- 35 labeled anti-TCR antibodies and subjected to flow cytometric analysis. CD8 +ve, TCR+ve T-lymphocytes were assessed for the number of cells possessing T-cell receptors capable of specifically recognizing and binding to OVA in the context of MHC Class I molecules.

[00210] For the cytotoxicity assay, the ability of splenocytes from immunized animals to specifically recognize and lyse target cells in an antigen specific manner was assessed using a 4 hour 51Chromium-release assay. Target cells were labeled with 51Chromium and the amount of cytotoxicity was determined by the amount of radionuclide released into the

5 supernatant from targets lysed by immune effector cells. Isolated splenocytes from immunized animals were tested immediately or after 5 days of in vitro restimulation with OVA-pulsed, syngeneic antigen presenting cells, for their ability to specifically lyse EG.7, OVA expressing target cells compared to EL4, non OVA-expressing cells.

10 [00211] The aim of the cytokine release assay is to detect antigen-specific immune effector cells that are activated to produce and secrete cytokines, specifically IFN- γ , in response to stimulation with a specific antigen. Cells were isolated from the spleen, liver, blood and lymph nodes of immunized animals and analyzed using the Cytokine Secretion Assay (Miltenyi Biotec). Cells were stimulated overnight with OVA-pulsed, autologous antigen presenting cells

15 and labeled with a catch reagent (a bispecific antibody recognizing the CD45 epitope on the surface of immune cells and IFN- γ). Any cells capable of recognizing and responding to the antigen stimulation, synthesized and secreted cytokines which were then captured by the cell-bound catch reagent, resulting in IFN- γ bound markers on their surface. Cells were then labeled with fluorescently labeled antibodies against IFN- γ and various phenotype markers and

20 analyzed by flow cytometry to allow detection of specific cell types that were activated to secrete IFN- γ .

[00212] Analysis of humoral response was designed to determine the level of antigen-specific IgG in the plasma of immunized mice. Blood was collected by cardiac puncture and centrifuged to collect plasma. Antigen specific immunoglobulin production was measured using the End-point dilution ELISA method to measure titers of total IgM, IgG and the IgG1, IgG2a, 25 subclasses. Samples of pooled plasma were serially diluted and plated into OVA coated plates to capture OVA specific antibodies in the diluted samples. OVA specific antibodies were then detected with horseradish peroxidase-conjugated rabbit anti-mouse IgM, IgG, IgG1, or IgG2a antibodies and TMB substrate. The absorbance of the colorimetric reaction was measured at 450nm on ELISA plate reader and end-point dilution titers were defined as highest dilution of plasma that resulted in absorbance value two times greater than that of naïve animals, with a cut-off value of 0.05. This was used to evaluate seroconversion and magnitude of response as well as to evaluate the Th type of response.

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35 [00213] Each of Figures 10 and 11 illustrate the normalization of ODN1m to that of its unmethylated counterpart ODN1. Each of the bars on these figures represents a direct comparison of one animal group (5 animals per group) treated with a methylated ODN and a second group treated with the unmethylated counterpart wherein each oligonucleotide is lipid

encapsulated in identical lipid particles. The results for the unmethylated population were set equivalent to 100% for each group and the methylated group was measured against this 100% standard. On bars showing an equivalence to 200%, this was the cut off value and in actuality the 200% line represents a value of 200% or greater.

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[00214] Figure 10 shows the results of the cytokine release assay described above. This figure illustrates that over a series of screenings, although both the methylated and unmethylated lipid encapsulated oligonucleotides each exhibited an immune response, on comparison of the methylated ODN to the unmethylated ODN, the methylated oligonucleotide 10 was as good as, and often better than, the unmethylated ODN in stimulating proliferation of dendritic cells, NK cells, and CD8+ T-cells as indicated by cytokine secretion in Figure 10A, B, and C respectively.

[00215] The results of the tetramer and CTL analyses are shown in Figures 11A - C. These 15 figures again illustrates the ability of both methylated and unmethylated ODN to stimulate an immune response. However, Figures 11A and B further demonstrate that over a series of screenings of animals treated with methylated or unmethylated encapsulated ODN, in each of the tetramer and CTL analyses respectively, the methylated oligonucleotide were consistently better in stimulating proliferation of cytotoxic T lymphocytes and tetrameric lymphocytes cells 20 than the unmethylated ODN. In addition, Figure 11C illustrates data from a representative tetramer study, wherein overall averages are shown in Figure 11B. Each of ODNS, ODN5m, ODN7 and ODN7m were tested as per the protocol described above. It is clearly shown in Figure 11C that lipid encapsulated ODN5m and ODN7m induce a higher number of antigen specific CD8 T-cells on comparison to their lipid encapsulated unmethylated counterparts.

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[00216] From each of Figures 10 and 11 it is shown that immune stimulation resulting from immunization with methylated ODN1m is consistently at least equivalent to, and often better than, the same treatment with its unmethylated oligonucleotide counterpart. This is further demonstrated in the following example.

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EXAMPLE 6

Prophylactic Anti-Tumor Efficacy Comparison of Methylated and Unmethylated Oligonucleotides in an EG7-OVA Tumor Model

[00217] The cancer vaccines provided herein include lipid-nucleic acid formulations in 35 conjunction with a tumor-associated antigen to stimulate an immune response to the antigen and the tumor *in vivo*. Hen egg albumin (ovalbumin; OVA) is a widely studied model antigen system. The antigenic determinants have been mapped and reagents and models exist to monitor both humoral and cell-mediated immune responses. In addition, cell lines containing

the OVA gene have been established and characterized and have been used routinely to evaluate anti-tumor immune responses following vaccination. Specifically, E.G7-OVA is a murine thymoma cell line engineered to express the OVA protein as a xenogeneic tumor-associated antigen and is an accepted model for investigating the factors required to induce a host's immune system to specifically attack malignant cells *in vivo*. A vigorous Th1 cytokine response and the induction of antigen-specific CD8+ T lymphocytes are considered essential for mounting an effective anti-tumor immune reaction.

[00218] Anti-tumor efficacy induced by subcutaneous immunization was assessed in C57Bl/6 (5 animals per group) in a prophylactic immunization model. Mice were immunized subcutaneously with 3 injections on a q7d x 3 dosing regimen on Days 0, 7 and 14 at a dose of 100 μ g oligonucleotide and 20 μ g of OVA antigen dose. Animals were then challenged with a subcutaneous injection of 2.5×10^6 EG.7 Ova expressing tumor cells on Day 21. Mice were monitored 3 times weekly to assess tumor growth and weight gain. Control mice were injected with one of PBS or HBS and 20 μ g of OVA antigen on the same schedule described above. Oligonucleotides were encapsulated in a lipid particle having a lipid composition of one of POPC: CHOL: DODAP:PEGCer14 or DSPC: CHOL: DODAP:PEGCer14 each in a ratio of 25:45:20:10. All comparisons of methylated and unmethylated oligonucleotides were done using like lipid particles. Results from these efficacy experiments are detailed in Figures 12 - 15, 18 - 21, and 25 - 28. Day 0 on each of the Figures is the day each animal was challenged with the tumor.

[00219] Figure 12 illustrates the efficacy trend when animals are immunized with free ODN. The results shown are consistent with the prior art, namely that when an animal is administered free oligonucleotides, the methylated oligonucleotides have less therapeutic efficacy than the unmethylated oligonucleotides in reducing tumor growth. Specifically, free unmethylated ODN1 and ODN2, having PS backbones so as to avoid nuclease degradation, showed a greater reduction in tumor growth than their methylated counterparts, ODN1m and ODN2m. This was most especially true about 25 days after inoculation with the tumor when the tumor growth rate of the methylated oligonucleotides approached the rate of the control animal treated only with a PBS buffer.

[00220] Figures 13-15 illustrate that encapsulation of oligonucleotides provides equivalent or better therapeutic efficacy of methylated over unmethylated oligonucleotides particularly when the oligonucleotides contain a natural phosphodiester (PO) backbone. Figure 13 shows that after implantation with a tumor, treatment with the methylated encapsulated ODN1m having a PS backbone was equal in therapeutic efficacy in comparison to the unmethylated ODN1. In contrast, Figure 14 shows the effect with the corresponding encapsulated methylated ODN1m and unmethylated ODN1 oligonucleotides having a PO backbone, where therapeutic efficacy

was greatest with the methylated version 32 days after transplantation while the unmethylated version lost its efficacy. Figure 15 shows that unmethylated ODN2 and its methylated counterpart ODN2m had virtually identical efficacy in reducing tumor growth. Accordingly, in certain embodiments the methylated oligonucleotide is at least as efficacious as an unmethylated counterpart when configured with a PS backbone.

5 [00221] Each of Figures 18, 19, 20 and 21 further elaborate on the above efficacy data. Figure 18 shows that lipid encapsulation of methylated PS-ODN1m provided a therapeutic benefit that was more effective than encapsulation of the PS-ODN1 in reducing tumor growth over a 10 prolonged period of time. This effectiveness was further borne out by the superior survival rates of mice treated with encapsulated PS-ODN1m in comparison to treatment with the PS-ODN1 in two different studies depicted in Figure 19. Figure 19A illustrates the percentage of animals that are tumor free at a series of time points and 19B, the number of animals remaining in the study at these same time points. As is clearly shown in Figure 19B, the number of 15 animals remaining in the study treated with ODN1 and ODN1m was essentially identical throughout the study. However, Figure 19A clearly illustrates a greater percentage of tumor free animals when treated with the methylated ODN1m compared to those treated with unmethylated ODN1.

20 [00222] Similarly, Figure 20 illustrates the tumor volume in mice treated with the two oligonucleotides over time and Figure 21 the percentage of animals surviving over time. Figure 20 shows improved efficacy when animals were treated with the encapsulated methylated ODN1m in comparison to the encapsulated unmethylated counterpart, ODN1. Correspondingly, Figure 21 shows an increase in the survival rate of mice treated with the 25 methylated ODN1m relative to treatment with unmethylated ODN1.

30 [00223] A further study efficacy study was conducted on the same tumor model using a different immunization protocol. In this study anti-tumor efficacy induced by subcutaneous immunization was assessed in C57Bl/6 (5 animals per group) in a prophylactic immunization model. Mice were immunized subcutaneously with 2 injections on a q7d x 2 dosing regimen on Days 0 and 7 at a dose of 100 μ g oligonucleotide and 20 μ g of antigen. Animals were then challenged with a subcutaneous injection of 5 x 10⁵ EG.7 Ova expressing tumor cells on Day 21. Mice were monitored 3 times weekly to assess tumor growth and weight gain. Control mice were injected with PBS on the same schedule described above. Oligonucleotides in 35 Figure 24(b) were encapsulated in a lipid particle having a lipid composition of DSPC: CHOL: DODAP:PEGCer14 each in a ratio of 25:45:20:10. All comparisons of methylated and unmethylated oligos were done using like lipid particles. Results from these efficacy experiments are detailed in Figure 24. Day 0 on the Figure is the day each animal was challenged with the tumor.

[00224] Figure 24 illustrates an example of treating the experimental tumor E-G7 using the lipid encapsulated PS-ODN1, PS-ODN2, each unmethylated, PS-ODN1m, methylated, in conjunction with an E-G7 OVA tumor antigen, which in this case was associated with the lipid particle by being attached to the surface thereof. Figure 24A shows that when the oligonucleotides were administered in the absence of the immunostimulatory lipid particle, the methylated PS-ODN1m had little effect on tumor growth. The corresponding unmethylated oligonucleotide PS-ODN1 was effective in reducing tumor volume while the unmethylated oligonucleotide PS-ODN2 was partially effective. Figure 24B shows that not only did encapsulation of the oligonucleotides in the lipid particle increase the effectiveness of the unmethylated PS-ODN2 to a level similar to ODN1, but also that the encapsulated methylated oligonucleotide PS-ODN1m was more effective than either of the encapsulated unmethylated oligonucleotides.

10 [00225] Figures 25-28 further illustrate that encapsulation of oligonucleotides provides equivalent or greater therapeutic efficacy for encapsulated methylated over unmethylated oligonucleotides. Figure 25 illustrates that lipid encapsulation of methylated PS-ODN5m provided a more effective therapeutic benefit than encapsulation of the equivalent unmethylated PS-ODN5 in reducing tumor growth over time. The effectiveness was further borne out by the superior survival rate of mice treated with encapsulated methylated PS-ODN5m in comparison to treatment with the unmethylated PS-ODN5 as shown in Figure 26. Figure 27 illustrates that while free unmethylated PS-ODN7 provides some anti-tumor benefit, free unmethylated PS-ODN 7 and PO-ODN7 as well as free methylated PS-ODN7 and PO-ODN7 were relatively ineffective in reducing tumor growth. However, lipid encapsulation of methylated PO-ODN7m provided effective therapeutic benefit in reducing tumor growth. Similarly, these trends were also illustrated in Figure 28 in the survival rate of mice treated with these same ODN.

15 [00226] Figures 25-28 further illustrate that encapsulation of oligonucleotides provides equivalent or greater therapeutic efficacy for encapsulated methylated over unmethylated oligonucleotides. Figure 25 illustrates that lipid encapsulation of methylated PS-ODN5m provided a more effective therapeutic benefit than encapsulation of the equivalent unmethylated PS-ODN5 in reducing tumor growth over time. The effectiveness was further borne out by the superior survival rate of mice treated with encapsulated methylated PS-ODN5m in comparison to treatment with the unmethylated PS-ODN5 as shown in Figure 26. Figure 27 illustrates that while free unmethylated PS-ODN7 provides some anti-tumor benefit, free unmethylated PS-ODN 7 and PO-ODN7 as well as free methylated PS-ODN7 and PO-ODN7 were relatively ineffective in reducing tumor growth. However, lipid encapsulation of methylated PO-ODN7m provided effective therapeutic benefit in reducing tumor growth. Similarly, these trends were also illustrated in Figure 28 in the survival rate of mice treated with these same ODN.

20 [00227] Figures 25-28 further illustrate that encapsulation of oligonucleotides provides equivalent or greater therapeutic efficacy for encapsulated methylated over unmethylated oligonucleotides. Figure 25 illustrates that lipid encapsulation of methylated PS-ODN5m provided a more effective therapeutic benefit than encapsulation of the equivalent unmethylated PS-ODN5 in reducing tumor growth over time. The effectiveness was further borne out by the superior survival rate of mice treated with encapsulated methylated PS-ODN5m in comparison to treatment with the unmethylated PS-ODN5 as shown in Figure 26. Figure 27 illustrates that while free unmethylated PS-ODN7 provides some anti-tumor benefit, free unmethylated PS-ODN 7 and PO-ODN7 as well as free methylated PS-ODN7 and PO-ODN7 were relatively ineffective in reducing tumor growth. However, lipid encapsulation of methylated PO-ODN7m provided effective therapeutic benefit in reducing tumor growth. Similarly, these trends were also illustrated in Figure 28 in the survival rate of mice treated with these same ODN.

25 [00228] Figures 25-28 further illustrate that encapsulation of oligonucleotides provides equivalent or greater therapeutic efficacy for encapsulated methylated over unmethylated oligonucleotides. Figure 25 illustrates that lipid encapsulation of methylated PS-ODN5m provided a more effective therapeutic benefit than encapsulation of the equivalent unmethylated PS-ODN5 in reducing tumor growth over time. The effectiveness was further borne out by the superior survival rate of mice treated with encapsulated methylated PS-ODN5m in comparison to treatment with the unmethylated PS-ODN5 as shown in Figure 26. Figure 27 illustrates that while free unmethylated PS-ODN7 provides some anti-tumor benefit, free unmethylated PS-ODN 7 and PO-ODN7 as well as free methylated PS-ODN7 and PO-ODN7 were relatively ineffective in reducing tumor growth. However, lipid encapsulation of methylated PO-ODN7m provided effective therapeutic benefit in reducing tumor growth. Similarly, these trends were also illustrated in Figure 28 in the survival rate of mice treated with these same ODN.

30 [00229] Figures 25-28 further illustrate that encapsulation of oligonucleotides provides equivalent or greater therapeutic efficacy for encapsulated methylated over unmethylated oligonucleotides. Figure 25 illustrates that lipid encapsulation of methylated PS-ODN5m provided a more effective therapeutic benefit than encapsulation of the equivalent unmethylated PS-ODN5 in reducing tumor growth over time. The effectiveness was further borne out by the superior survival rate of mice treated with encapsulated methylated PS-ODN5m in comparison to treatment with the unmethylated PS-ODN5 as shown in Figure 26. Figure 27 illustrates that while free unmethylated PS-ODN7 provides some anti-tumor benefit, free unmethylated PS-ODN 7 and PO-ODN7 as well as free methylated PS-ODN7 and PO-ODN7 were relatively ineffective in reducing tumor growth. However, lipid encapsulation of methylated PO-ODN7m provided effective therapeutic benefit in reducing tumor growth. Similarly, these trends were also illustrated in Figure 28 in the survival rate of mice treated with these same ODN.

EXAMPLE 7

Therapeutic Anti-Tumor Efficacy Comparison of Methylated and Unmethylated

Oligonucleotides In a B-16 Melanoma Tumor Model

[00226] Anti-tumor efficacy induced by intravenous tail immunization was assessed in C57Bl/6 (8 animals per group) in a therapeutic immunization model. Animals were challenged with a subcutaneous injection of 3.0×10^5 EG.7 B16/BL6 murine melanoma expressing tumor cells on Day-0. Mice were then treated intravenously every other day starting on day 4 for 14 days at a dose of 20mg/kg ODN. Mice were monitored every other day to assess tumor growth and weight gain. Control mice were injected with HBS on the same schedule described above. Oligonucleotides were encapsulated in a lipid particle having a lipid composition of DSPC:CHOL: DODAP:PEGCer14 each in a ratio of 25:45:20:10. Results from this efficacy

experiments are detailed in Figures 16 and 17. Day 0 on each of the Figures is the day each animal was challenged with the tumor.

[00227] Figure 16 illustrates therapeutic efficacy of administering the methylated PS-ODN1m to 5 an animal inoculated with a B16 melanoma tumor in comparison to its unmethylated counterpart PS-ODN1. Encapsulation of PS-ODN1m in a lipid particle increased its efficacy in reducing tumor volume to at least that of the encapsulated unmethylated PS-ODN1.

[00228] Figure 17 illustrates the average weight of the tumors in each mouse on Day 22. The 10 average tumor size in mice treated with free methylated PS-ODN1m was nearly the same as in mice treated with a buffer control, while mice treated with free unmethylated PS-ODN1 showed reduced tumor growth. In contrast, when mice were treated with the methylated PS-ODN1 encapsulated in a lipid particle, the amount of tumor reduction was near equivalent to that obtained with the lipid encapsulated unmethylated PS-ODN1. Accordingly, lipid encapsulation 15 of methylated oligonucleotides can yield efficacy in treating a tumor *in vivo* even though the free methylated oligonucleotide has little or no efficacy.

EXAMPLE 8

Blood Clearance Levels when Treated with Encapsulated Oligonucleotides

20 [00229] An important aspect in effective immune stimulation is the ability of the immune system to raise an antibody response against specific antigens. One of the first demonstrations of the capacity of antigen associated with lipid encapsulated oligonucleotides to initiate such a response is illustrated by the data shown in Figure 23.

25 [00230] Each of ODN1 and ODN1m were encapsulated in two different lipid particles; Lipid one (L1) being a DSPE:CHOL:DODAP:PEGCer20 and the Lipid 2 (L2) being the same but having a PEGCer14 in the place of the PEGCer20. The half-life of the PEGCer 20 within the liposome is known to be much longer than that of the PEGCer14 and thus the PEGCer20 30 remains with the lipid particle for a longer time period. Mice were given a series of 4 i.v. tail injections, starting on Day 0, and were dosed once a week for 3 weeks. Blood was collected 1-hour post injection each week and were analyzed for the presence of the encapsulated ODN.

35 [00231] Figure 23 illustrates the effect on clearance from the blood in mice for the different lipid compositions, L1 and L2 each with different PEG-ceramide steric coatings (PEG-ceramide-C-20 and PEG-ceramide C-14 respectively) in combination with either methylated or unmethylated oligonucleotides, ODN1m and ODN1 respectively. After injection 1 the results show extended circulation/slow clearance for both of the encapsulated ODNs from the blood sample regardless of composition. However, for each of injections 2, 3 and 4 the results show that L1 liposomes

(L1-ODN1 and L1-ODN1m) containing the long-lived PEGCer20 had shorter circulation/rapid clearance while L2-ODN1 and L2-ODN1m containing the short-lived PEG-ceramide C-14 had longer circulation/slower clearance than those encapsulated with lipid particles containing PEGCer20 lipid.

5

[00232] The data depicted herein demonstrates two specific points: (1) the induction of antigen specific antibodies; and (2) the relative immunostimulatory capacity of unmethylated and methylated oligonucleotide. In terms of induction of antigen specific antibodies, the initial injection resulted in the induction of antibodies directed against the PEG moiety of the PEG-10 ceramide steric barrier lipid. The presence of these antibodies in the plasma of injected animals resulted in the opsonization and subsequent rapid clearance from the circulation of liposomes containing PEG after injections 2, 3 and 4 as seen with L1 liposomes with PEGCer20. However, animals injected with liposomes without PEG, such as L2 liposomes with PEGCer14 from which the PEG dissociated very rapidly in circulation, were not opsonized and 15 thus had relatively extended circulation times. In terms of the relative immunostimulatory potency of unmethylated vs. methylated oligonucleotide, the clearance of the liposomes containing either the unmethylated oligonucleotide or the corresponding methylated form were cleared at similar rates, thus indicating that both are able to induce antigen specific antibodies.

20 **EXAMPLE 9****Induction of CTL Response Using A Polytope Approach with Multiple Tumor-Associated Antigens**

[00233] Single epitope-based approaches have the disadvantage that an MHC-restricted CTL 25 response is raised to only one antigen. In addition many cancer antigens are non-mutated differentiation antigens, such as TRP-2 exemplified above, and thus self-reactive T cells in the host are predominantly deleted during thymic education. A polytope approach would allow multiple antigens to be simultaneously targeted and should increase the spectrum of anti-tumor CTL responses against such self-antigens. CTL responses specific for multiple antigens and 30 restricted by multiple MHC alleles would clearly be desirable for broader immune reactivity, given the variable expression of tumor antigens and MHC alleles in different malignancies. Targeting multiple antigens associated with a particular malignancy would minimize the chances of tumor escape by antigen downregulation or epitope mutation.

35 [00234] Targeting multiple antigens and MHC alleles might be achieved by using multiple recombinant antigen mixtures of synthetic peptide epitopes. To improve the immune response against these multi-epitope antigens several adjuvants are under assay. This experiment employs encapsulated PS-ODN 1m using POPC:Chol:DODMA:PEG-DMG in combination with two murine melanoma antigens TRP2 (H₂K_b, VYDFFVWL) and Gp100 (H₂D_b, EGSRNQDWL).

C57BL/6 mice were injected 3 times with either antigen, or both, in the presence of encapsulated PS-ODN 1m. As positive control we used dendritic cells known to be potent in inducing CTL responses. B16 lysate containing multiple epitopes was also assayed either with ODN 1m or DC for the ability to induce multi-epitope immunity. Immune response was 5 assessed by the ability of CD8+ T cells to mediate cytotoxicity against tumor cells in an antigen-specific manner.

Results: When injected together with encapsulated ODN 1m, a CTL response was raised against both antigens (Figure 29). PS-ODN 1m was as good as DC in generating CTL 10 response against both antigens delivered together (Figure 30). Injection of B16 lysate with PS-ODN 1m was more potent in inducing CTL than injection of DC incubated with B16 lysate (Figure 31).

[00235] The Examples provided illustrate certain embodiments of the invention. In a more 15 general sense, however, the invention encompasses compositions and methods for providing therapeutic benefits to mammalian subjects (including humans) utilizing such compositions. The compositions of the invention are in the form of a lipid membrane vesicle; and a nucleic acid fully encapsulated within said vesicle. Where stimulation of a response to a particular antigen is desired, the composition may also associate the antigen with the vesicle, for example 20 via chemical coupling, hydrophobic bonding or ionic bonding to an external surface of the vesicle, or encapsulation within the vesicle.

[00236] Preferred compositions are those in which the nucleic acid comprises greater than 4% by weight of the composition.

25 [00237] The nucleic acid in the compositions of the invention may suitably be nucleic acids which are not complementary to the genome of the treated mammal, and which provide immunostimulation through a mechanism which does not depend on a complementary base-pairing interaction with nucleic acids of the mammal. Such nucleic acids will frequently contain 30 an immunostimulating sequence, such as a CpG motif or an immune stimulating palindrome.

[00238] The nucleic acids used in the compositions of the invention may be nucleic acids which do not induce an immune response when administered in free form to a naïve mammal, or which suppress an immune response to an immune stimulating sequence of nucleotides when 35 administered in free form to a naïve mammal.

[00239] The nucleic acids may have exclusively phosphodiester internucleotide linkages or may be modified in which a way that they a plurality of phosphodiester internucleotide linkages in combination with modified internucleotide linkages. The nucleic acids may also contain

exclusively modified linkages, or a plurality of modified linkages. For example, the nucleic acid may contain exclusively phosphorothioate internucleotide linkages or a plurality of phosphorothioate internucleotide linkages.

- 5 [00240] The cationic lipid which is used in formulating the composition suitably is selected from DODAP, DODMA, DMDMA, DOTAP, DC-Chol, DDAB, DODAC, DMRIE, and DOSPA. In addition, the lipid formulation preferably includes an aggregation preventing compound, such as a PEG-lipid, a PAO-lipid or a ganglioside.
- 10 [00241] In addition to or instead of an antigen, the compositions of the invention can include a co-encapsulated cytotoxic agent such as doxorubicin. The lipid membrane vesicle fully encapsulates both the nucleic acid and the cytotoxic agent. Compositions of this type can be prepared by a method which is a further aspect of the invention. In this method, a therapeutic composition is prepared preparing lipid in ethanol; mixing lipid with oligonucleotide in aqueous buffer to form oligonucleotide loaded lipid vesicles; and exposing the oligonucleotide loaded lipid vesicles to a cytotoxic agent such that the cytotoxic agent actively accumulates in the interior space of said vesicle.
- 15 [00242] The compositions of the invention can be used in various methods to provide therapeutic benefits to mammals, including humans, through the use of a lipid-nucleic acid particle comprising a nucleic acid which is fully encapsulated in a lipid formulation comprising a cationic lipid in the manufacture of a medicament. Thus, the compositions can be used to induce an immune response in a mammal, to activate CTL or B cells in a mammal or to treat neoplasia in a mammal having a neoplasia by a method comprising the steps of preparing a lipid-nucleic acid particle comprising a nucleic acid which is fully encapsulated in a lipid formulation, which lipid formulation comprises a cationic lipid; and administering the lipid-nucleic acid particle to the mammal.
- 20 [00243] When an antigen is included in the composition, the invention provides a method of inducing an immune response to the antigen comprising preparing a particle comprising a lipid membrane vesicle comprising a nucleic acid fully encapsulated within said vesicle and an antigen to which an immune response is desired associated with an external surface of said vesicle, and administering the particles to the mammalian subject to be treated.
- 25 [00244] As demonstrated in the examples above, the utilization of a lipid carrier in the compositions in accordance with the invention allows a substantial reduction in the amount of oligonucleotide needed to achieve the desired stimulation of the immune system. In some cases, this is reflected in the fact that an oligonucleotide which had no apparent activity in the free form is useful for stimulating an immune response when provided in lipid-encapsulated

form. In other cases, this is reflected in the fact that the amount of ODN necessary to achieve the same level of response with a lower dosage of ODN. Thus, in practicing a method employing an effective amount of oligonucleotide to stimulate an immune response in a mammal, the present invention provides the improvement comprising fully-encapsulating the 5 oligonucleotide in a lipid vesicle and administering less than 20% of said effective amount of oligonucleotide to a mammalian subject, thereby obtaining a desired immune response in said mammalian subject.

[00245] While the data depicted herein demonstrates immunostimulatory activity *in vivo* and 10 therapeutic efficacy using certain exemplary embodiments of the invention, which are provided for completeness and consistency, it is understood that the invention is not limited to these exemplary embodiments. One of ordinary skill in the art will be readily able to make and use other specific embodiments of the invention consistent with the teachings provided herein.

CLAIMS

1. A lipid-methylated nucleic acid formulation for stimulating an immune response in an animal, said formulation comprising a lipid component and a nucleic acid component comprising a methylated nucleic acid sequence.
2. The lipid-nucleic acid formulation according to Claim 1, wherein said methylated nucleic acid sequence comprises at least one CpG dinucleotide having a methylated cytosine.
3. The formulation according to Claim 2, wherein said methylated cytosine comprises a methyl or hydroxymethyl group attached to the carbon-4 position (4-mC) or carbon-5 position (5-mC).
4. An adjuvant comprising a lipid-nucleic acid (LNA) formulation, wherein said LNA formulation comprises:
 - a) a lipid component comprising at least one cationic lipid; and
 - b) a nucleic acid component comprising at least one methylated oligonucleotide;wherein said adjuvant is capable of stimulating dendritic cells *in vivo* in response to antigenic stimulation.
5. The adjuvant according to Claim 4, wherein said adjuvant is capable of stimulating dendritic cell expansion *in vivo* characterized by an increase in the number of antigen-presenting cells expressing at least one of a CD11c and a DEC205 marker.
6. The adjuvant according to Claim 4, wherein said adjuvant is capable of stimulating dendritic cell activation *in vivo* characterized by an increase in the number of antigen-presenting cells co-expressing at least one of a CD11c and a DEC205 marker in conjunction with a CD86 marker.
7. The adjuvant according to Claim 4, wherein said at least one methylated oligonucleotide comprises a single CpG dinucleotide having a methylated cytosine.
8. The adjuvant according to Claim 7, wherein said at least one methylated oligonucleotide comprises the sequence 5' TAACGTTGAGGGGCAT 3' (ODN1m).
9. The adjuvant according to Claim 4, wherein said at least one methylated oligonucleotide comprises two CpG dinucleotides, and wherein at least one of the cytosines in said CpG dinucleotides is methylated.

10. The formulation according to Claim 9, wherein said methylated nucleic acid sequence is an oligonucleotide having the sequence 5' TTCCATGACGTTCTGACGTT 3' (ODN2m).
11. The adjuvant according to Claim 4, wherein said at least one methylated oligonucleotide comprises at least one CpG dinucleotide having a methylated cytosine.
12. The adjuvant according to any one of Claims 4 to 11, wherein said oligonucleotide comprises a modified phosphate backbone.
13. The adjuvant according to Claim 12, wherein said modified phosphate backbone is phosphorothioate.
14. A vaccine comprising a lipid-nucleic acid (LNA) formulation in combination with at least one target antigen, wherein said at least one target antigen is mixed with or associated with said LNA formulation, said formulation comprising:
 - a) a lipid component comprising at least one cationic lipid; and
 - b) a nucleic acid component comprising at least one methylated oligonucleotide, wherein said vaccine is capable of stimulating dendritic cells *in vivo* in response to presentation of said at least one target antigen by said formulation to antigen-presenting cells.
15. The vaccine according to Claim 14, wherein said vaccine is capable of stimulating dendritic cell expansion *in vivo* characterized by an increase in the number of antigen-presenting cells expressing at least one of a CD11c and a DEC205 marker.
16. The vaccine according to Claim 14, wherein said vaccine is capable of stimulating dendritic cell activation *in vivo* characterized by an increase in the number of antigen-presenting cells co-expressing at least one of a CD11c and a DEC205 marker in conjunction with a CD86 marker.
17. The vaccine according to Claim 14, wherein said at least one methylated oligonucleotide comprises a single CpG dinucleotide having a methylated cytosine.
18. The vaccine according to Claim 14, wherein said at least one methylated oligonucleotide comprises a plurality of CpG dinucleotides, and wherein at least one of said CpG dinucleotides comprises a methylated cytosine.
19. The vaccine according to Claim 14, wherein said at least one methylated oligonucleotide comprises at least one CpG dinucleotide having a methylated cytosine.

20. The vaccine according to any one of Claims 14 to 19, wherein said at least one target antigen comprises a microbial antigen.

21. The vaccine according to any one of Claims 14 to 19, wherein said at least one target antigen comprises a tumor-associated antigen.

22. The vaccine according to Claim 14, wherein said at least one target antigen comprises a plurality of epitopes from the same antigen.

23. The vaccine according to Claim 14, wherein said at least one target antigen comprises a plurality of epitopes from different antigens.

24. The vaccine according to any one of Claims 14 to 23, wherein said oligonucleotide comprises a modified phosphate backbone.

25. The vaccine according to Claim 24, wherein said modified phosphate backbone is phosphorothioate.

26. A method for stimulating an enhanced host immune response to antigenic stimulation comprising administering to said host a lipid-nucleic acid (LNA) formulation, said LNA formulation comprising:

- a) a lipid component comprising at least one cationic lipid; and
- b) a nucleic acid component comprising at least one methylated oligonucleotide;

wherein said LNA formulation is capable of stimulating dendritic cells *in vivo* in response to antigenic stimulation.

27. The method according to Claim 23, wherein said LNA formulation is capable of stimulating dendritic cell expansion *in vivo* characterized by an increase in the number of antigen-presenting cells expressing at least one of a CD11c and a DEC205 marker.

28. The method according to Claim 23, wherein said LNA formulation is capable of stimulating dendritic cell activation *in vivo* characterized by an increase in the number of antigen-presenting cells co-expressing at least one of a CD11c and a DEC205 marker in conjunction with a CD86 marker.

29. The method according to Claim 26, wherein said at least one methylated oligonucleotide comprises a single CpG dinucleotide having a methylated cytosine.

30. The method according to Claim 29, wherein said at least one methylated oligonucleotide comprises the sequence 5' TAACGTTGAGGGGCAT 3' (ODN1m).
31. The method according to Claim 26, wherein said at least one methylated oligonucleotide comprises at least two CpG dinucleotides, and wherein the cytosine in at least one of said CpG dinucleotides is methylated.
32. The method according to Claim 31, wherein said methylated nucleic acid sequence is an oligonucleotide having the sequence 5' TTCCATGACGTTCTGACGTT 3' (ODN2m).
33. The method according to any of Claims 26 to 32, wherein said LNA formulation is administered in combination with at least one target antigen, wherein said at least one target antigen is mixed with or associated with said LNA formulation.
34. The method according to Claim 33, wherein said at least one target antigen is mixed with said LNA formulation.
35. The method according to Claim 33, wherein said at least one target antigen is associated with said LNA formulation.
36. The method according to any one of Claims 26 to 35, wherein said methylated oligonucleotide comprises a modified phosphate backbone.
37. The method according to Claim 36, wherein said methylated oligonucleotide comprises a phosphorothioate backbone.
38. A method for stimulating dendritic cells, comprising contacting at least one dendritic cell with a lipid-methylated nucleic acid formulation comprising a lipid component and a nucleic acid component comprising a methylated nucleic acid sequence.
39. A method for stimulating host dendritic cells *in vivo*, comprising administering to said host a lipid-methylated nucleic acid formulation comprising a lipid component and a nucleic acid component comprising a methylated nucleic acid sequence, wherein said formulation is capable of stimulating dendritic cells *in vivo* in response to antigenic stimulation.
40. The method according to Claim 38 or 39, wherein stimulating dendritic cells comprises dendritic cell expansion characterized by an increase in the number of antigen-presenting cells expressing at least one of a CD11c and a DEC205 marker.

41. The method according to Claim 38 or 39, wherein stimulating dendritic cells comprises dendritic cell activation characterized by an increase in the number of antigen-presenting cells co-expressing at least one of a CD11c and a DEC205 marker in conjunction with a CD86 marker.

42. The method according to any one of Claims 38 to 41, wherein said methylated nucleic acid sequence comprises at least one CpG dinucleotide having a methylated cytosine.

43. The method according to Claim 42, wherein said methylated cytosine comprises a methyl or hydroxymethyl group attached to the carbon-4 position (4-mC) or carbon-5 position (5-mC).

44. A method for simultaneously delivering antigenic and adjuvant immune stimulation to antigen presenting cells, comprising the administration of a lipid-nucleic acid (LNA) formulation associated with a target antigen, said LNA formulation comprising:

- a) a lipid component comprising at least one cationic lipid; and
- b) a nucleic acid component comprising at least one oligonucleotide having at least one

CpG dinucleotide;

wherein said LNA formulation is capable of stimulating dendritic cells *in vivo*.

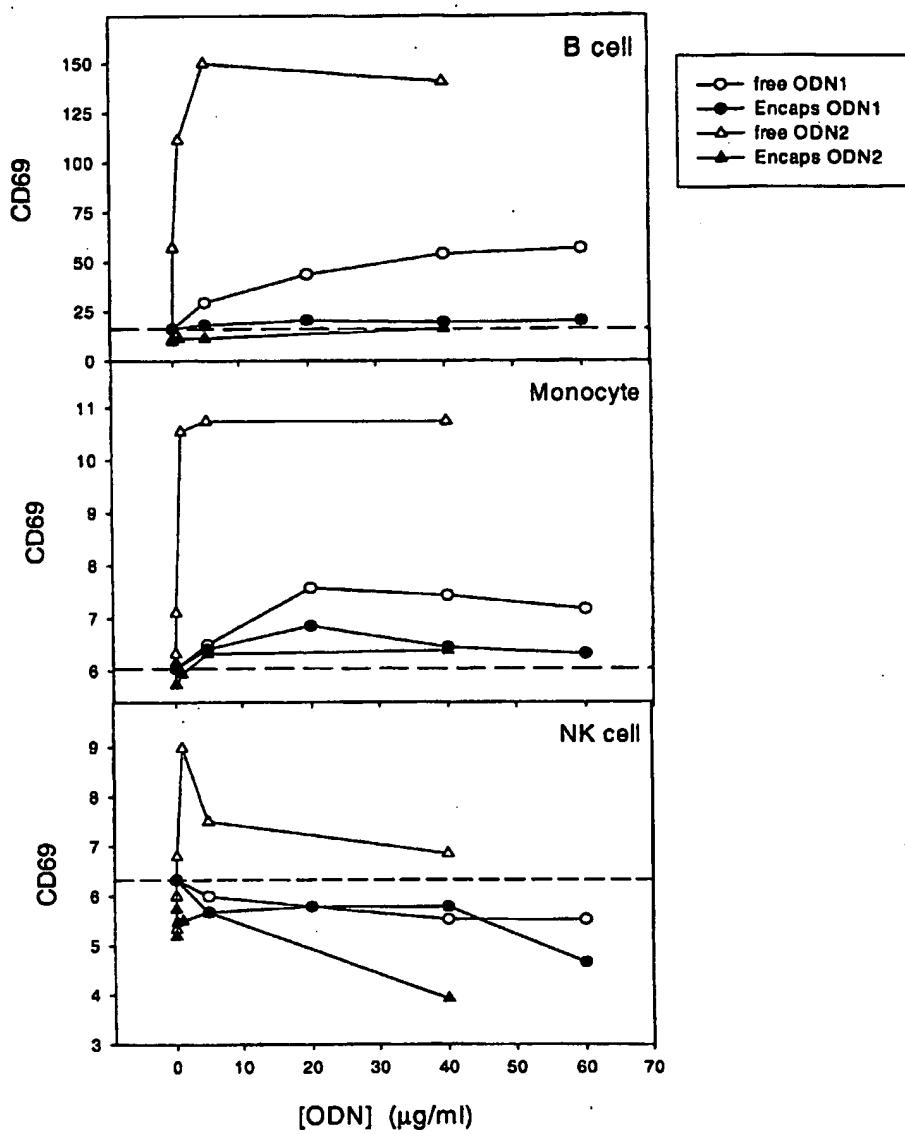


Figure 1

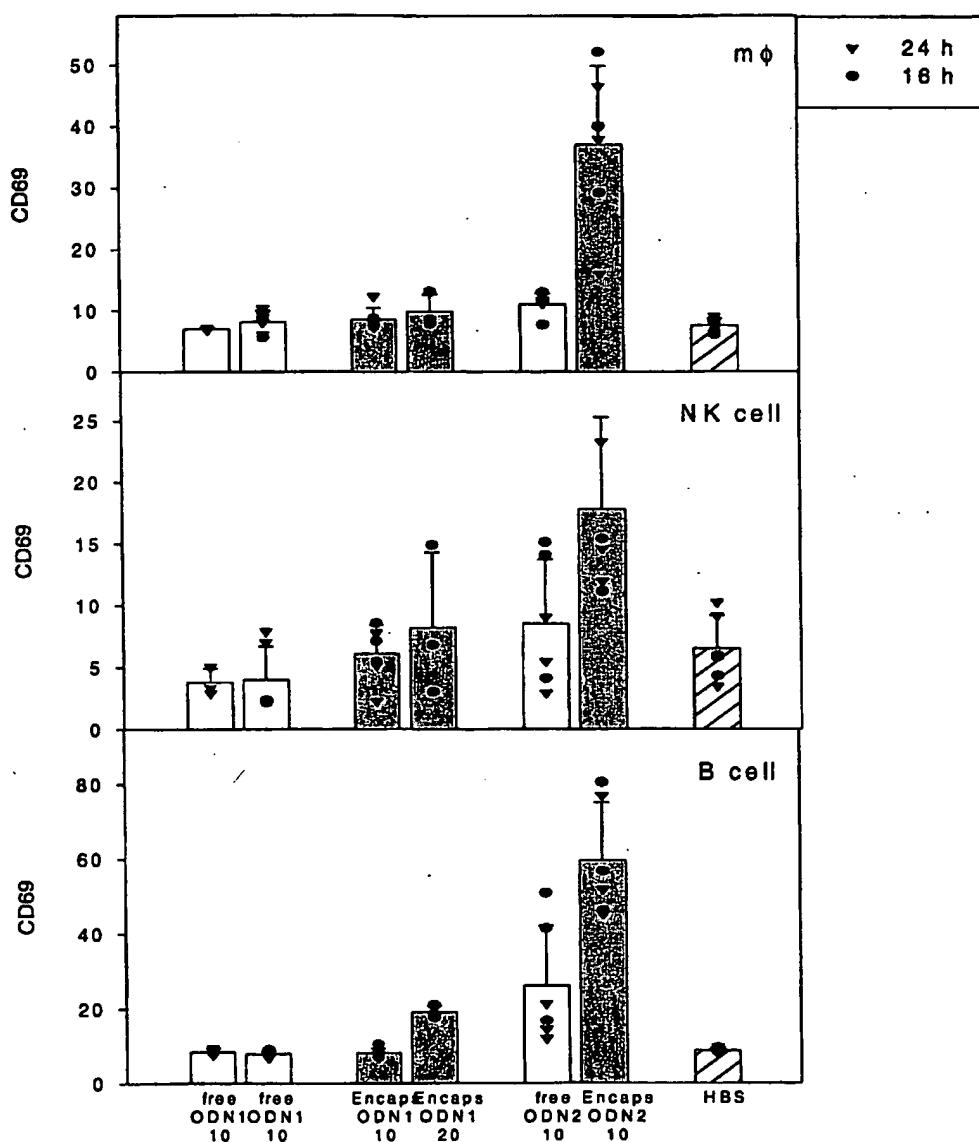


Figure 2

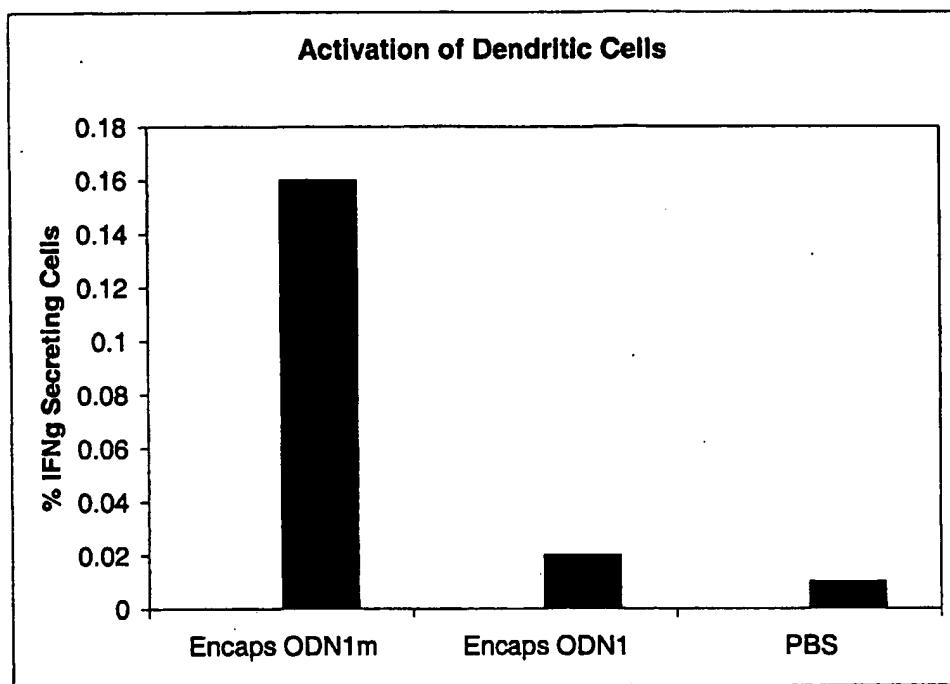


Figure 3

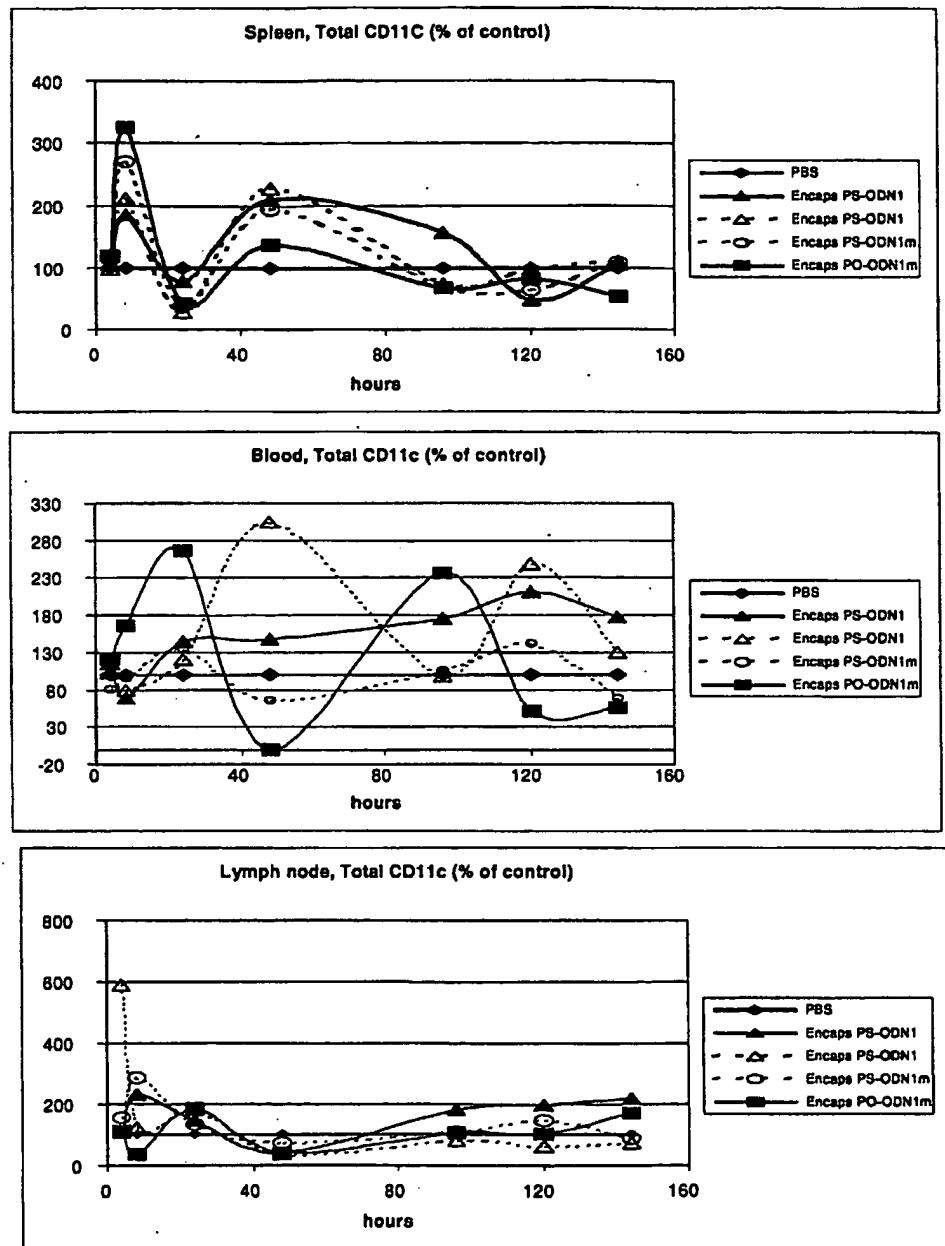


Figure 4A

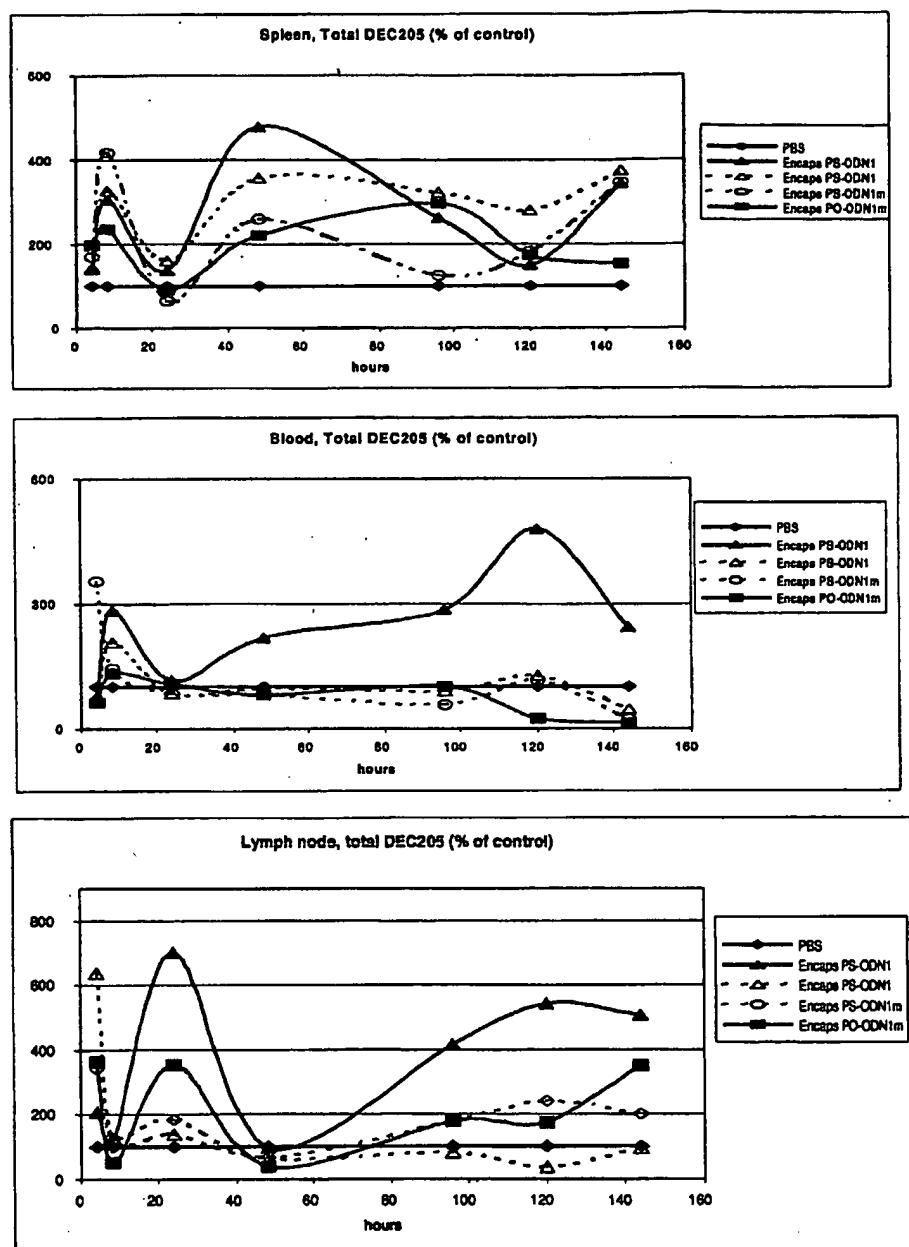


Figure 4B

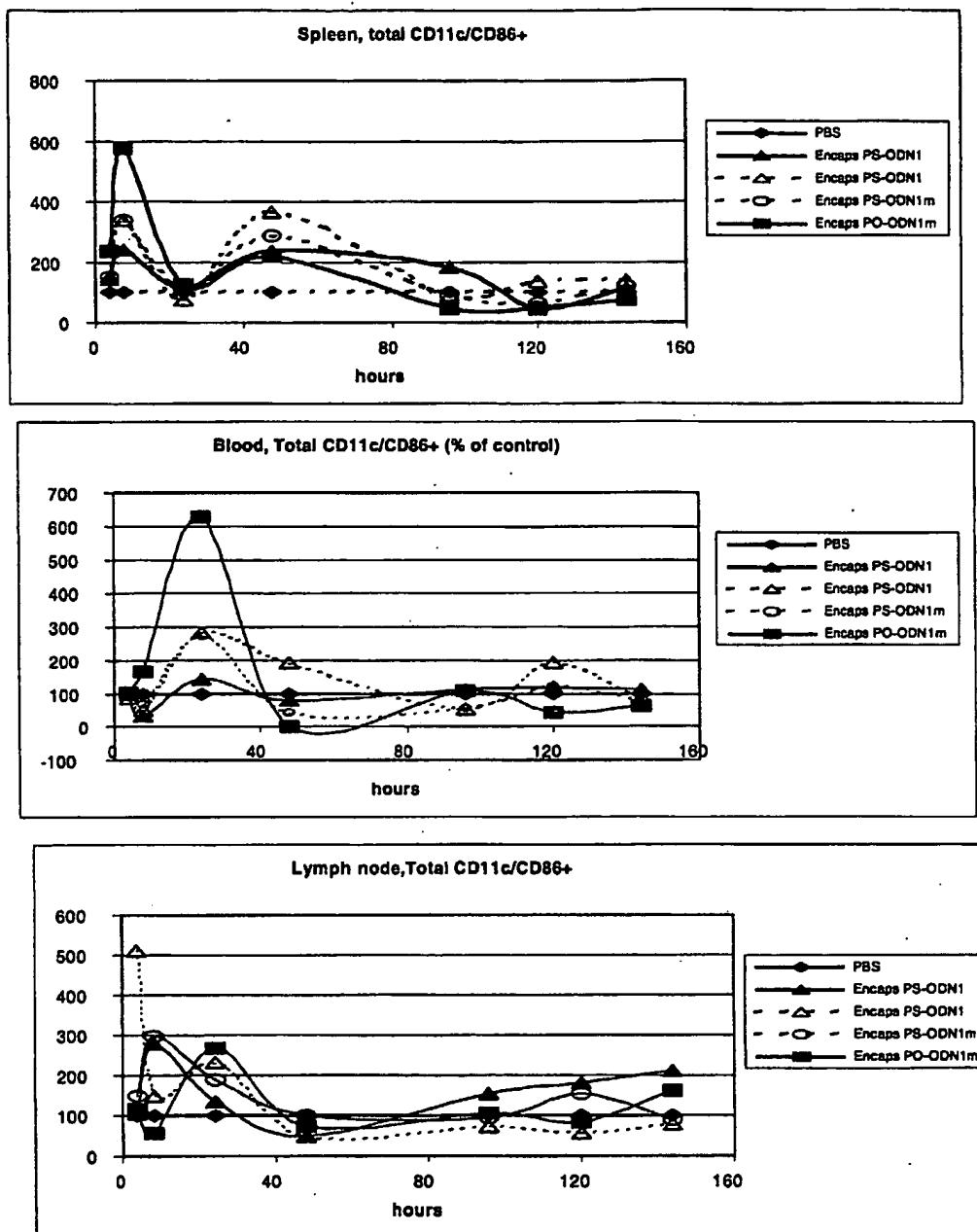


Figure 5A

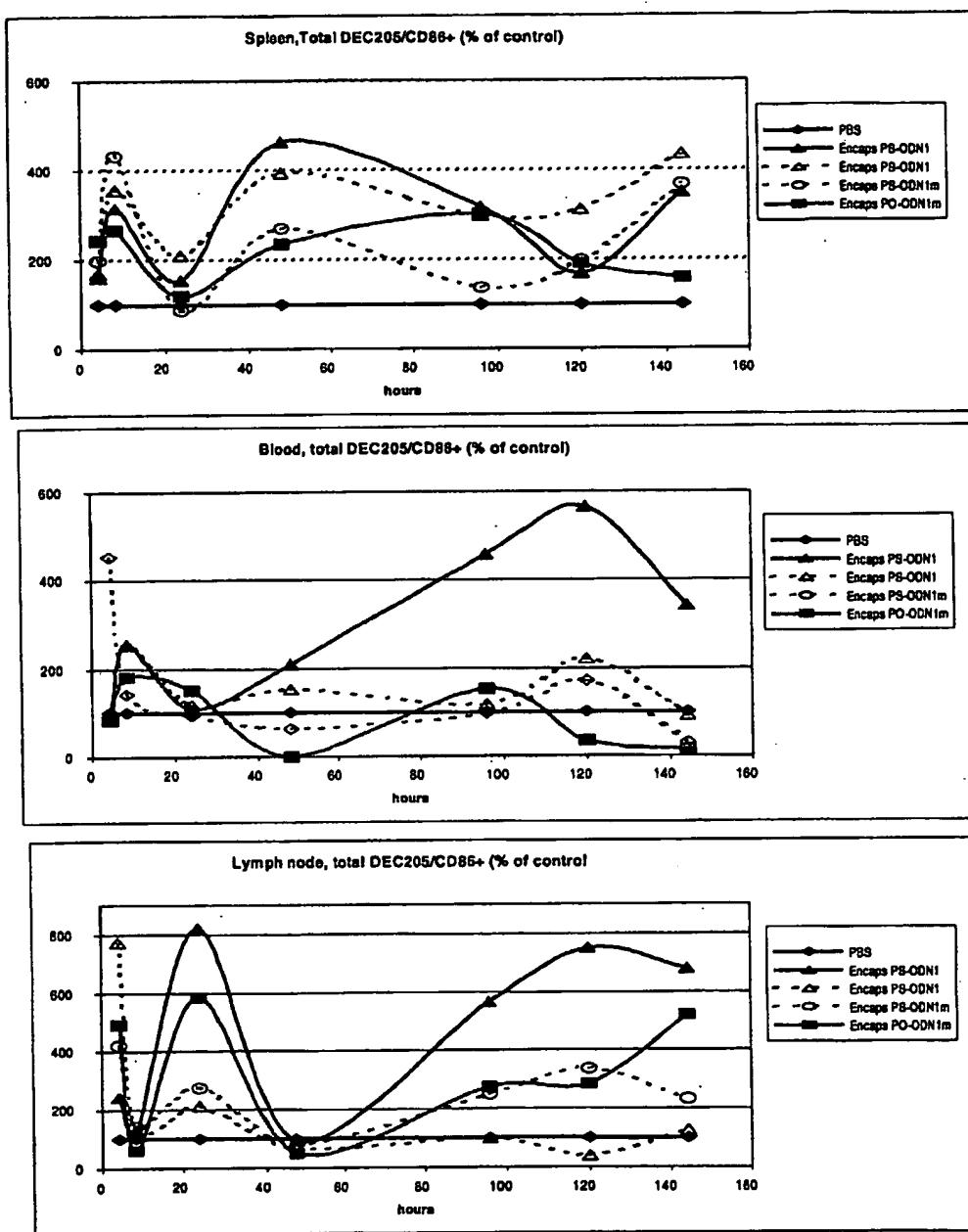


Figure 5B

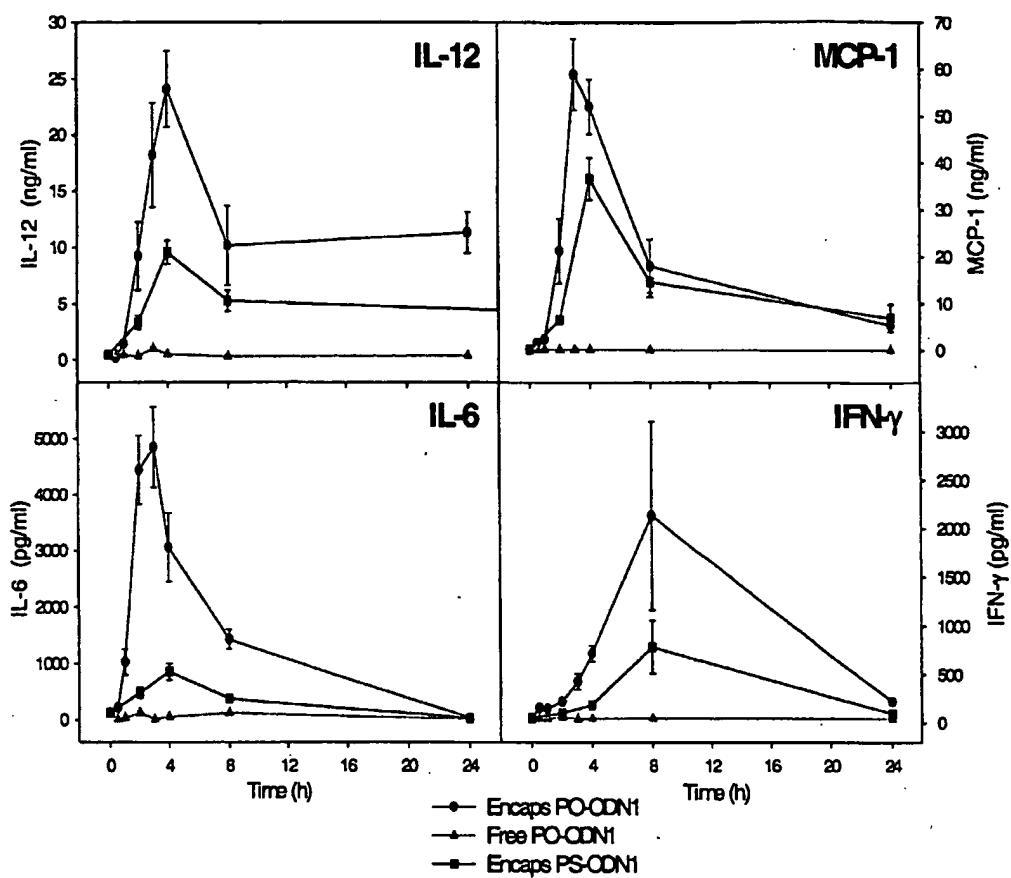


Figure 6

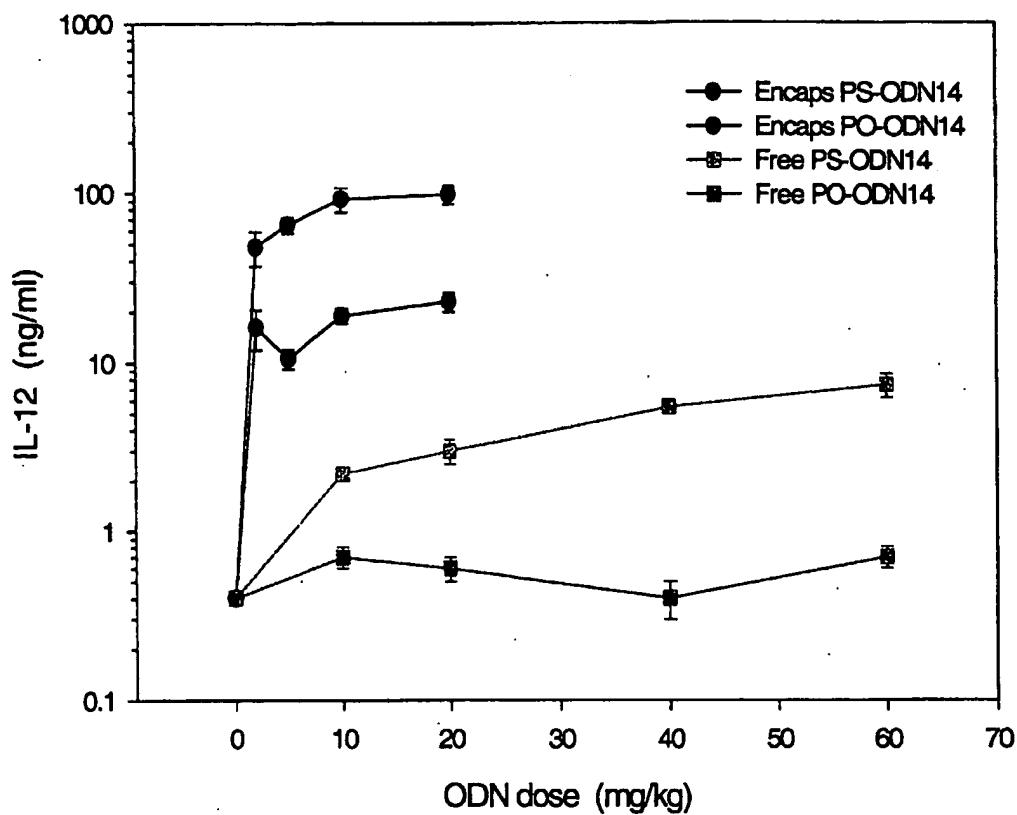


Figure 7A

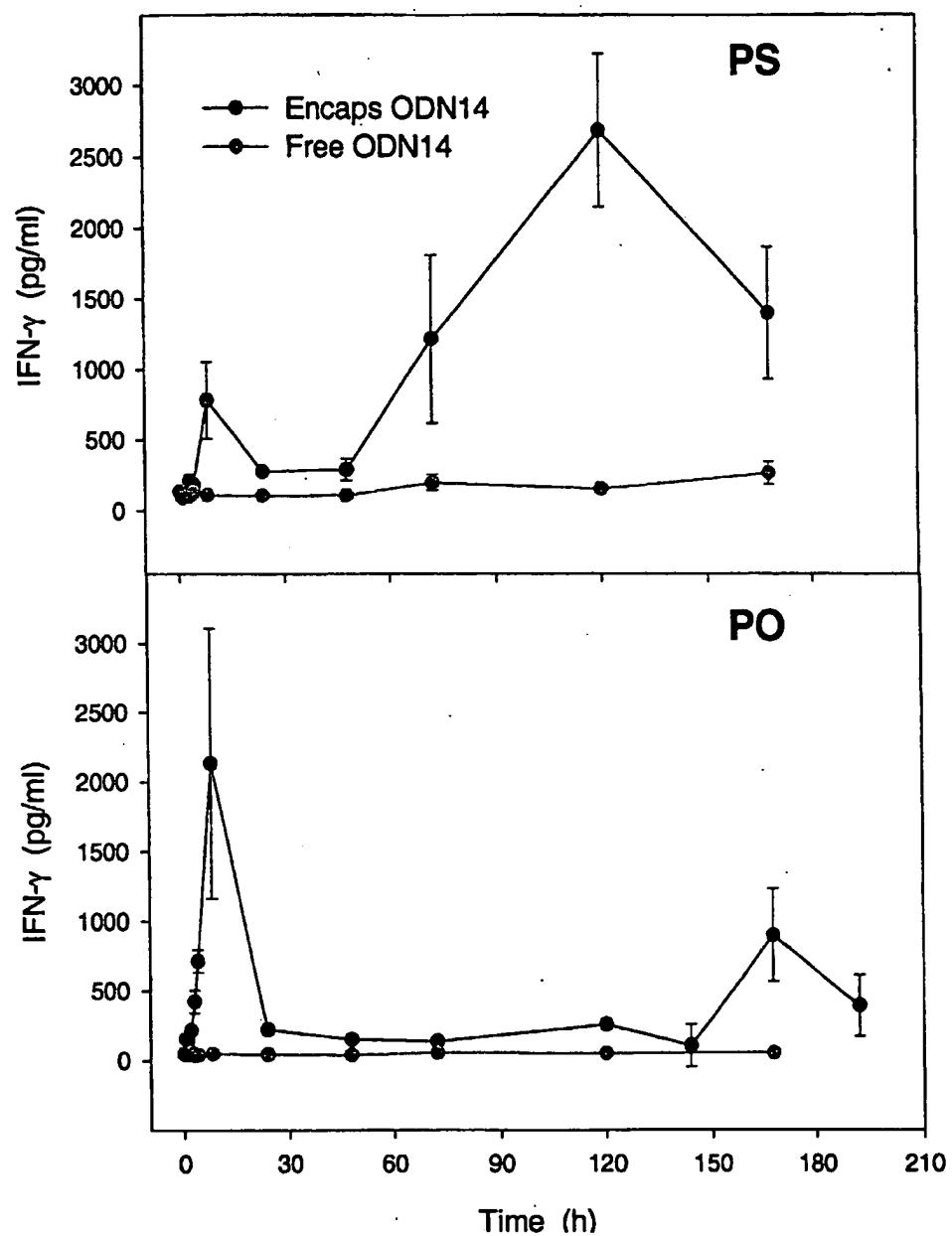


Figure 7B

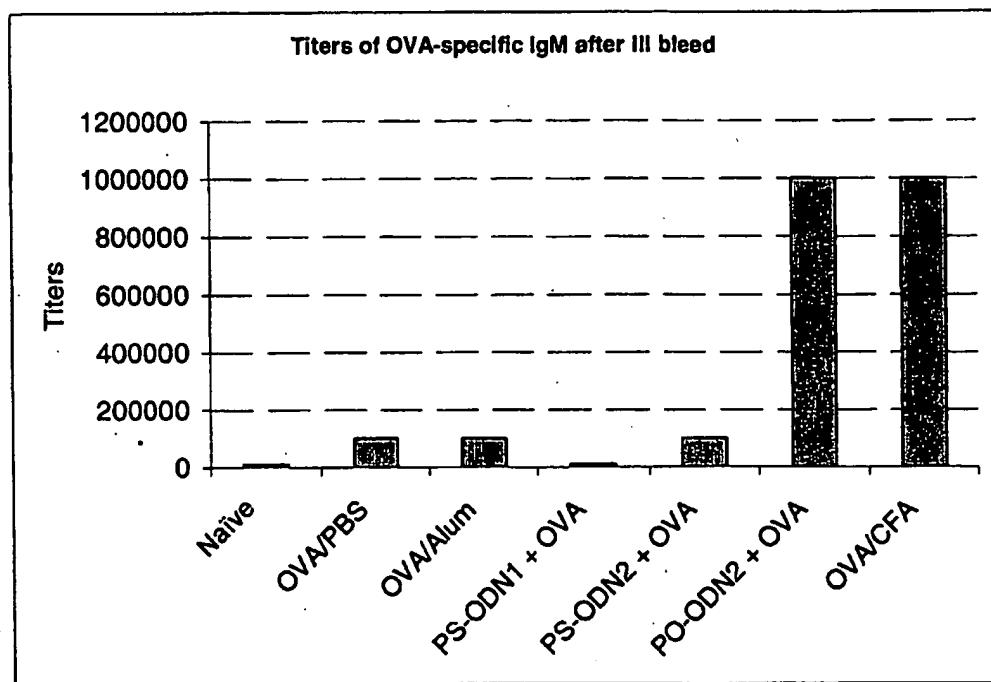


Figure 8

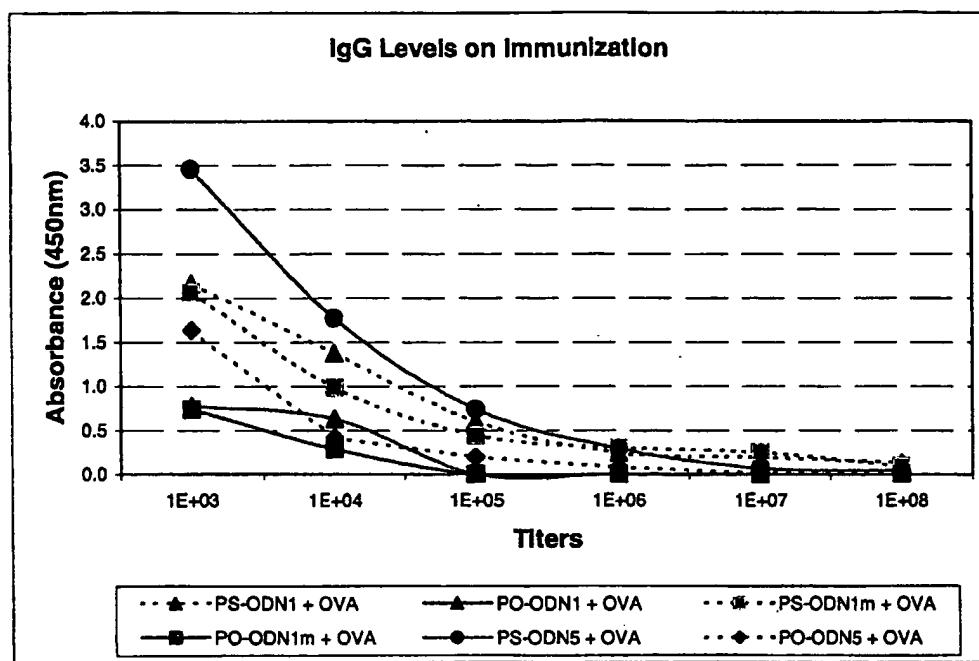


Figure 9

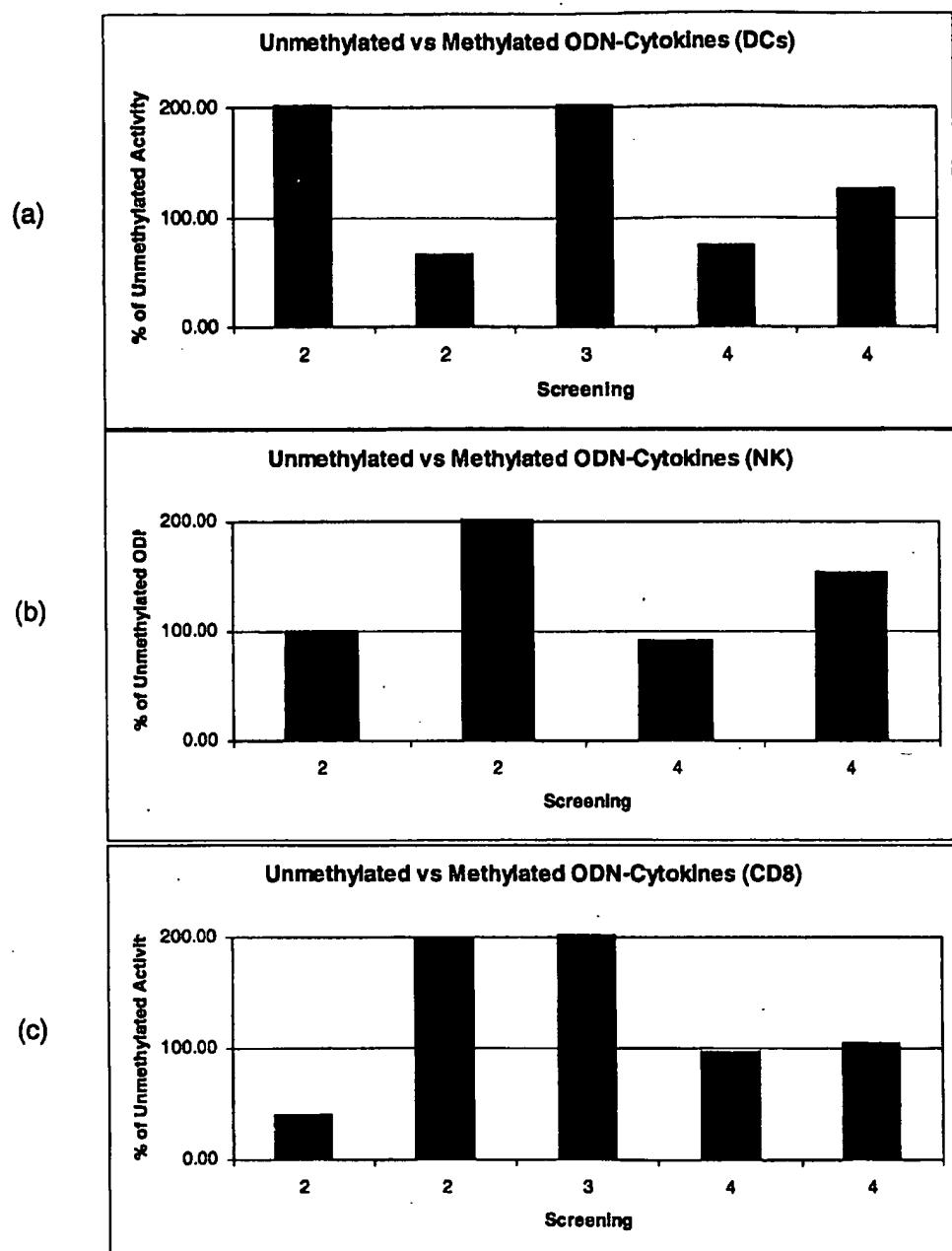


Figure 10

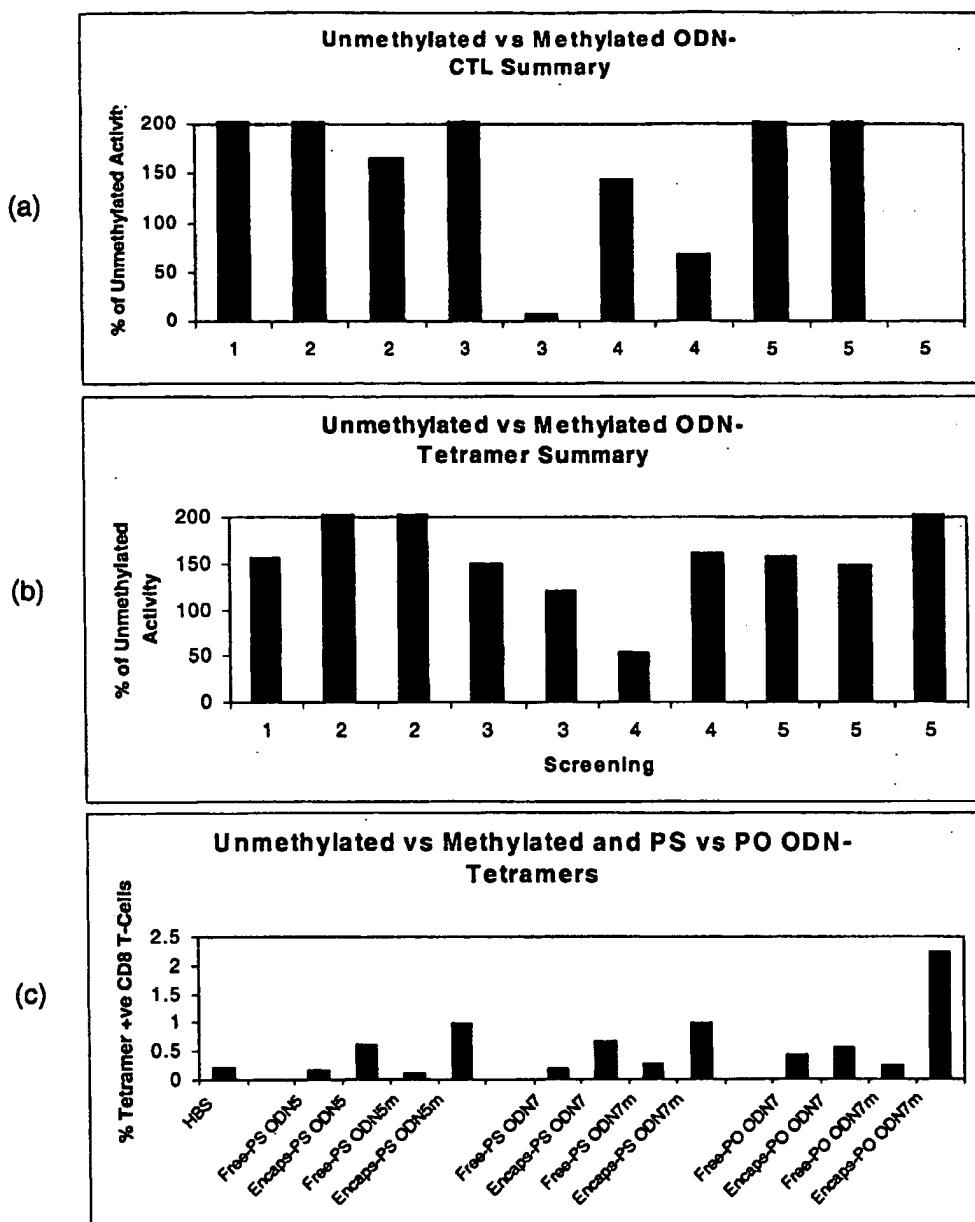


Figure 11

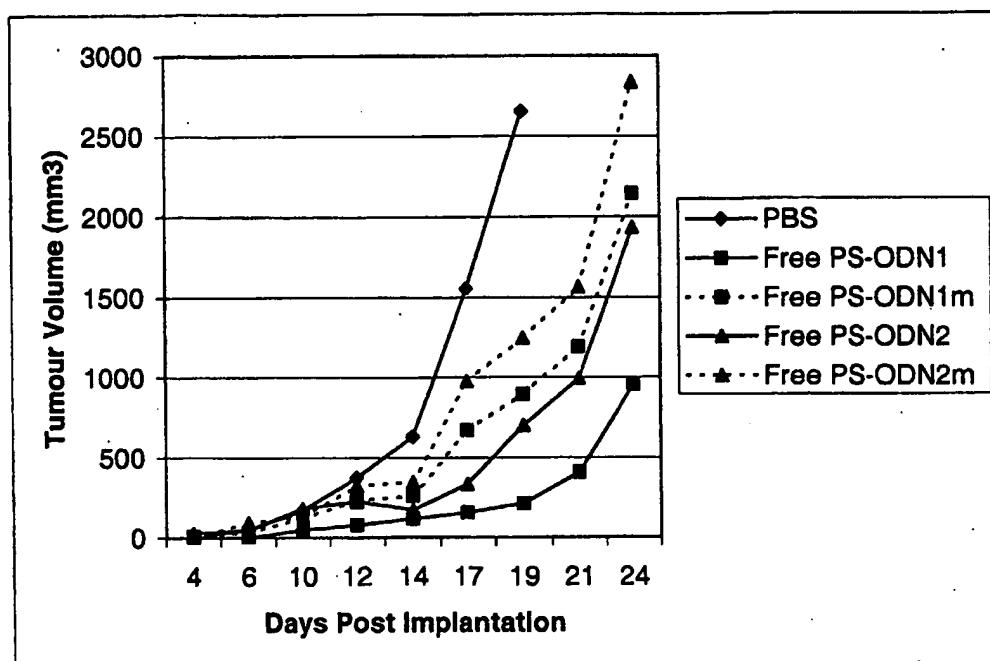


Figure 12

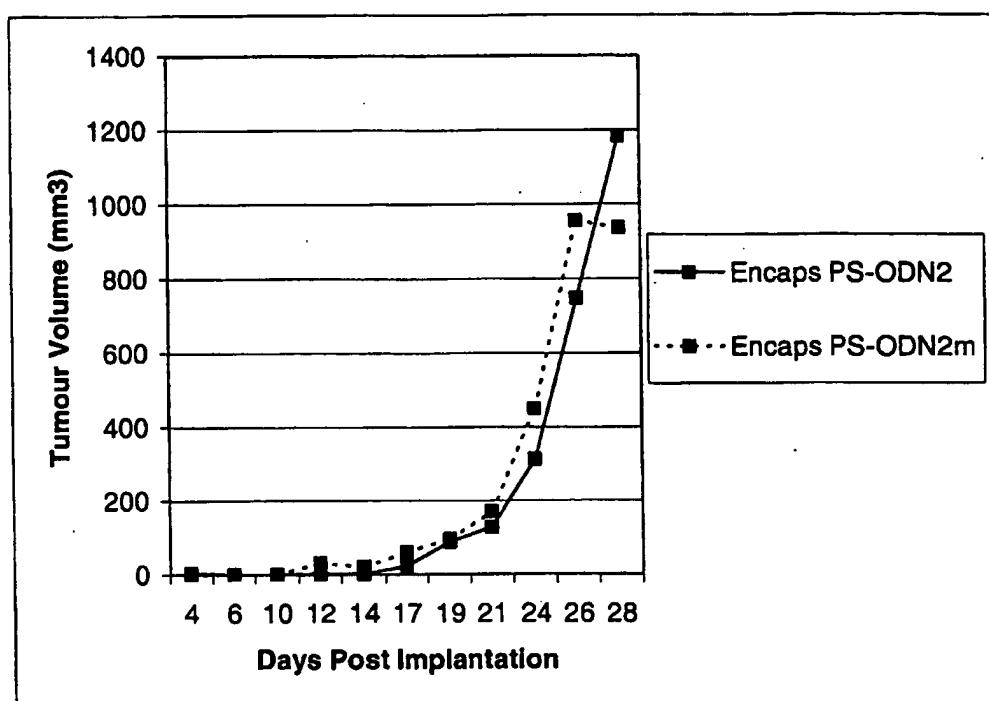


Figure 13

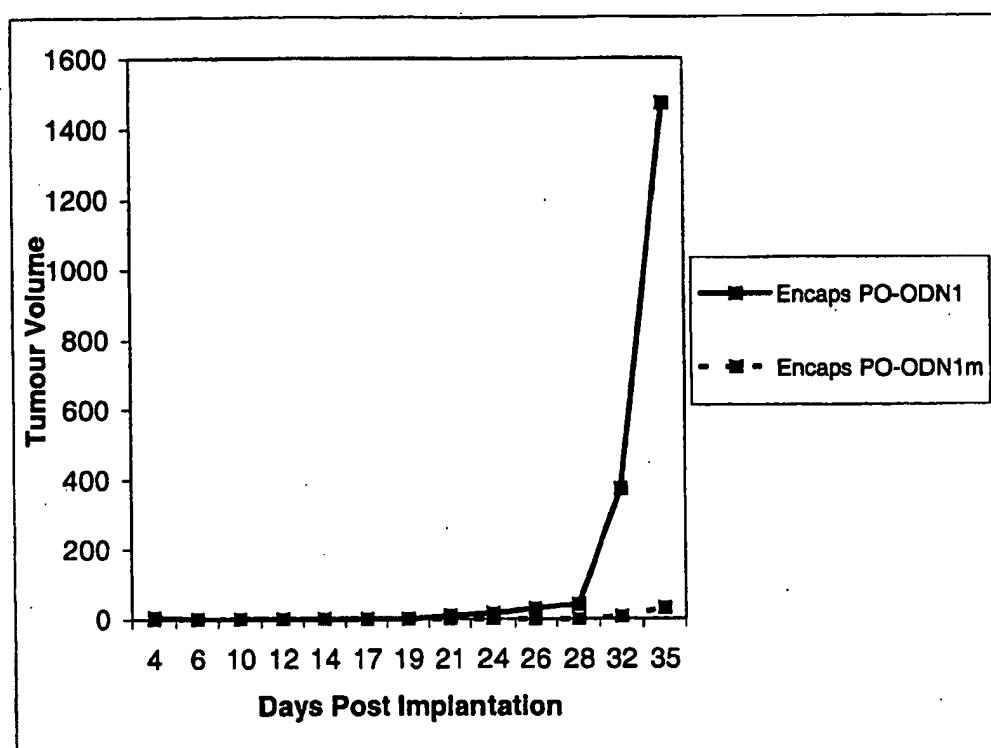


Figure 14

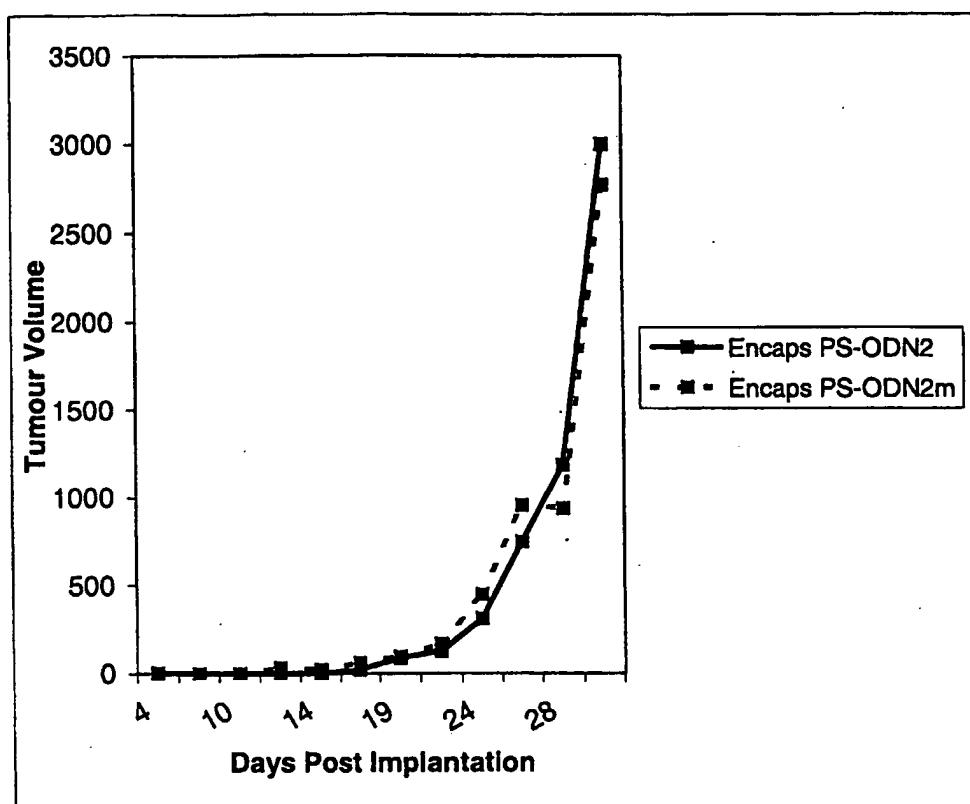


Figure 15

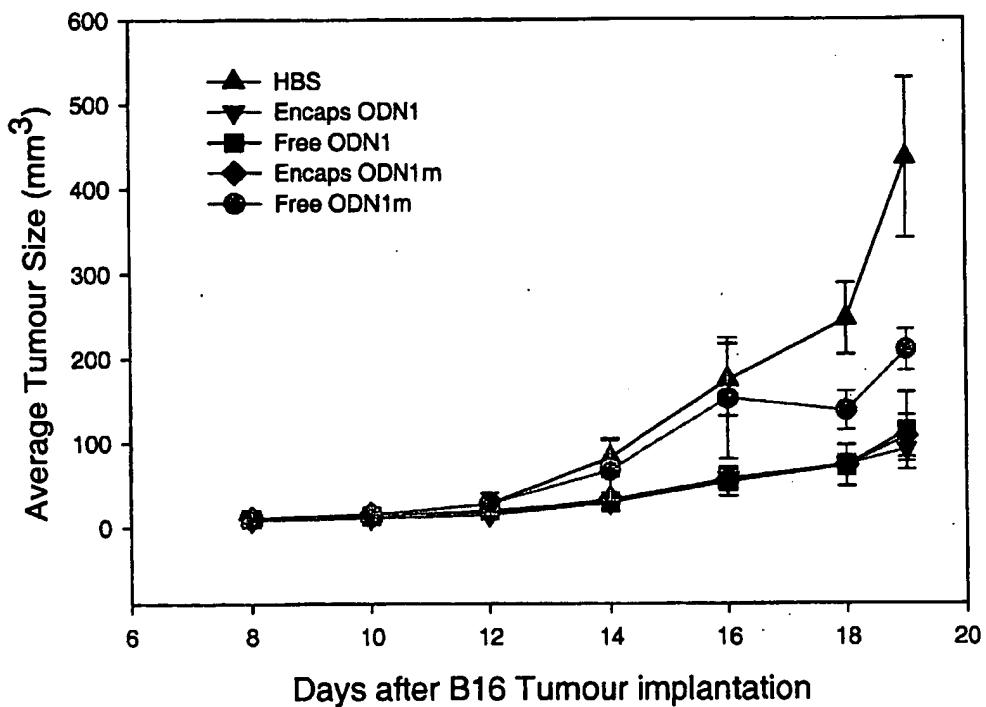


Figure 16

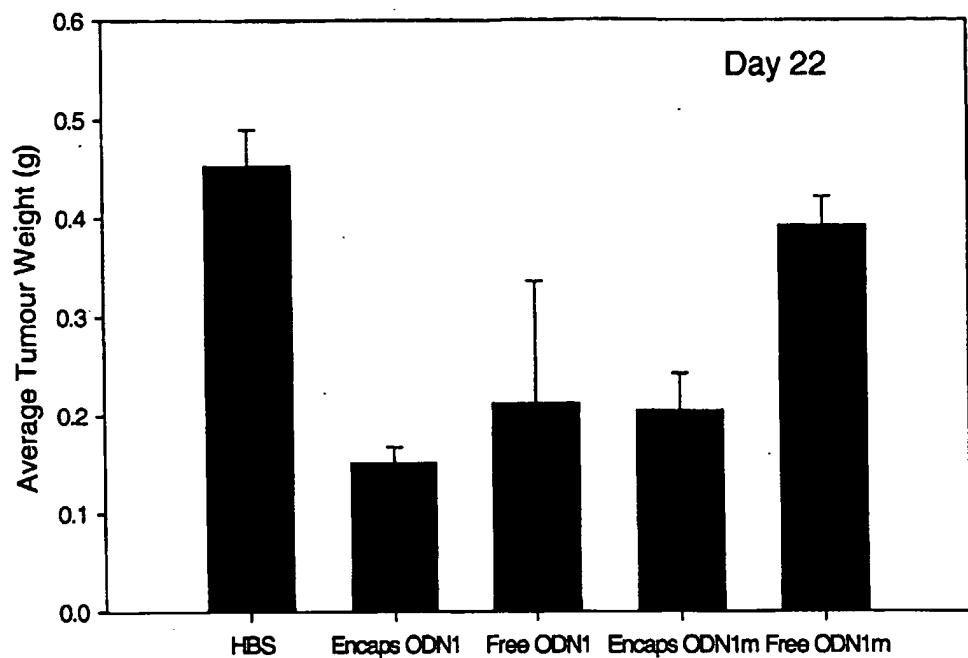


Figure 17

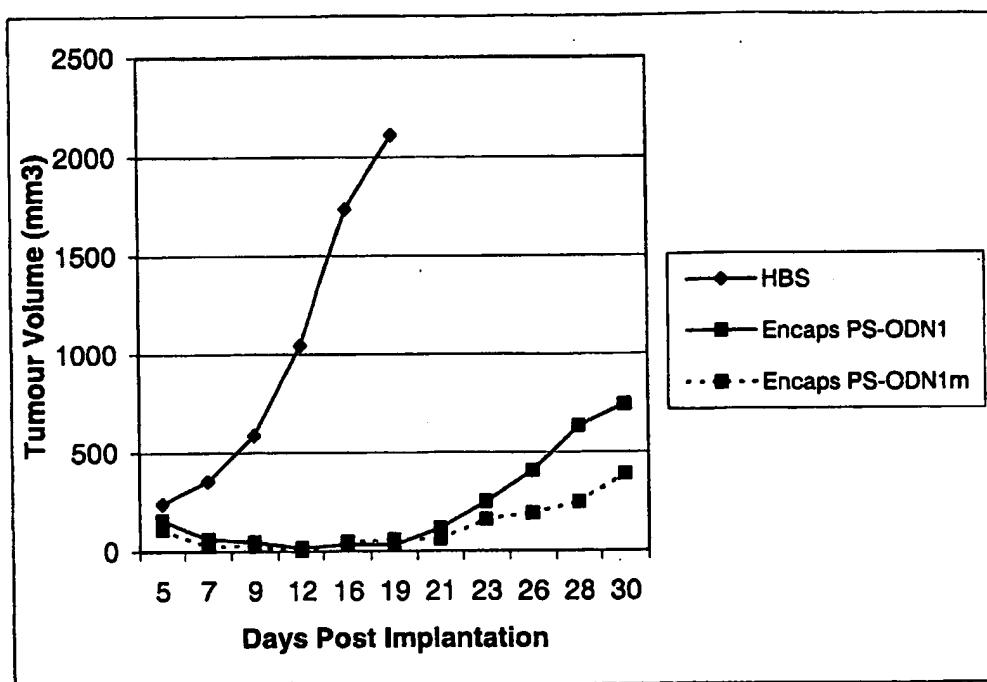


Figure 18

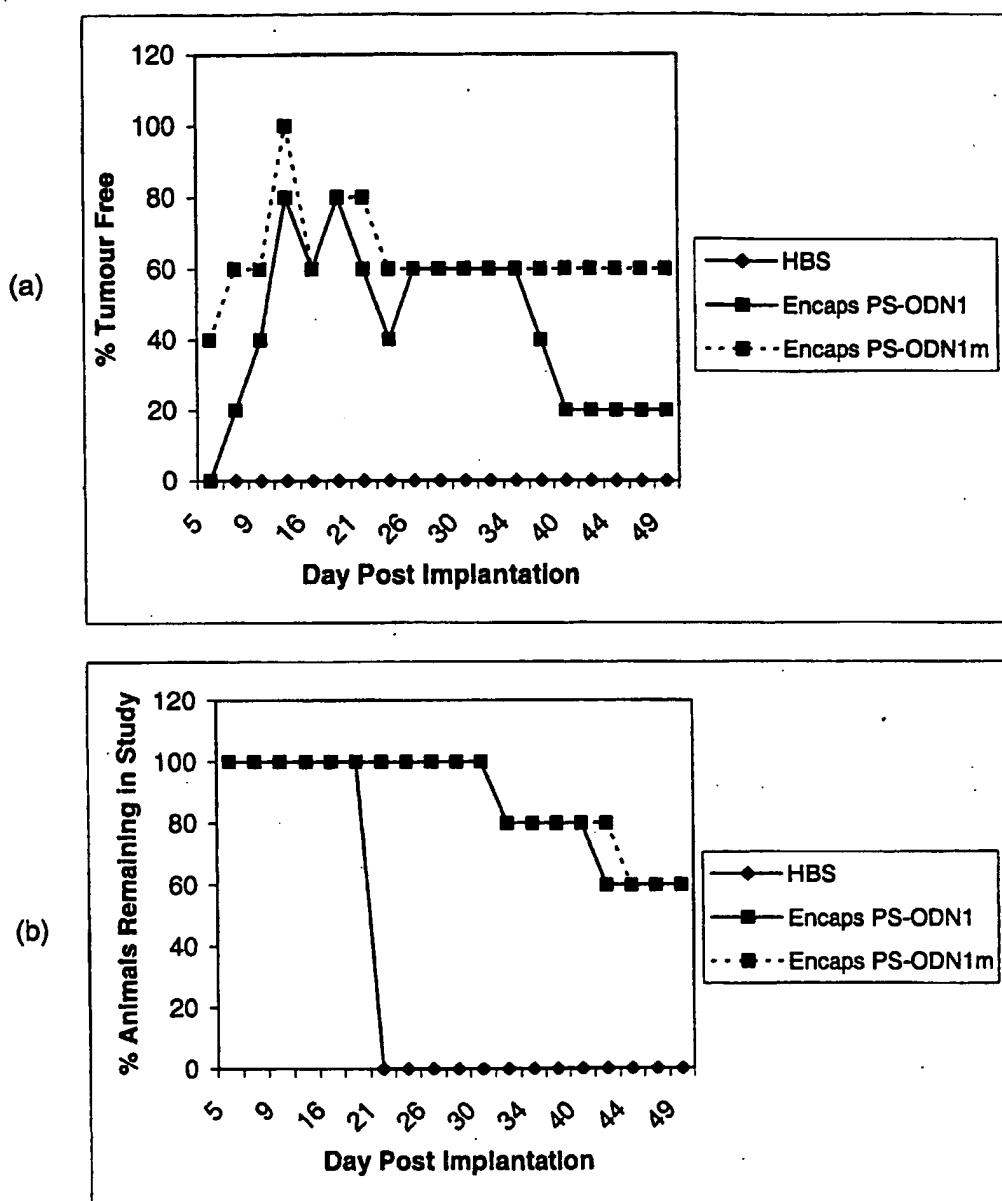


Figure 19

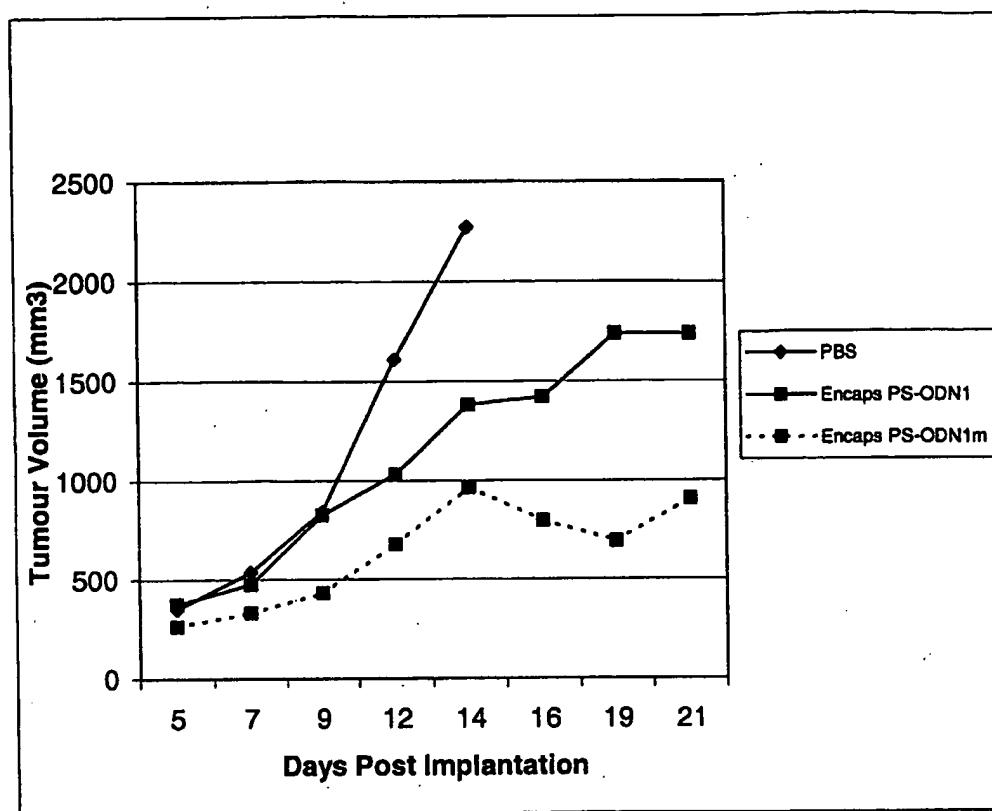


Figure 20

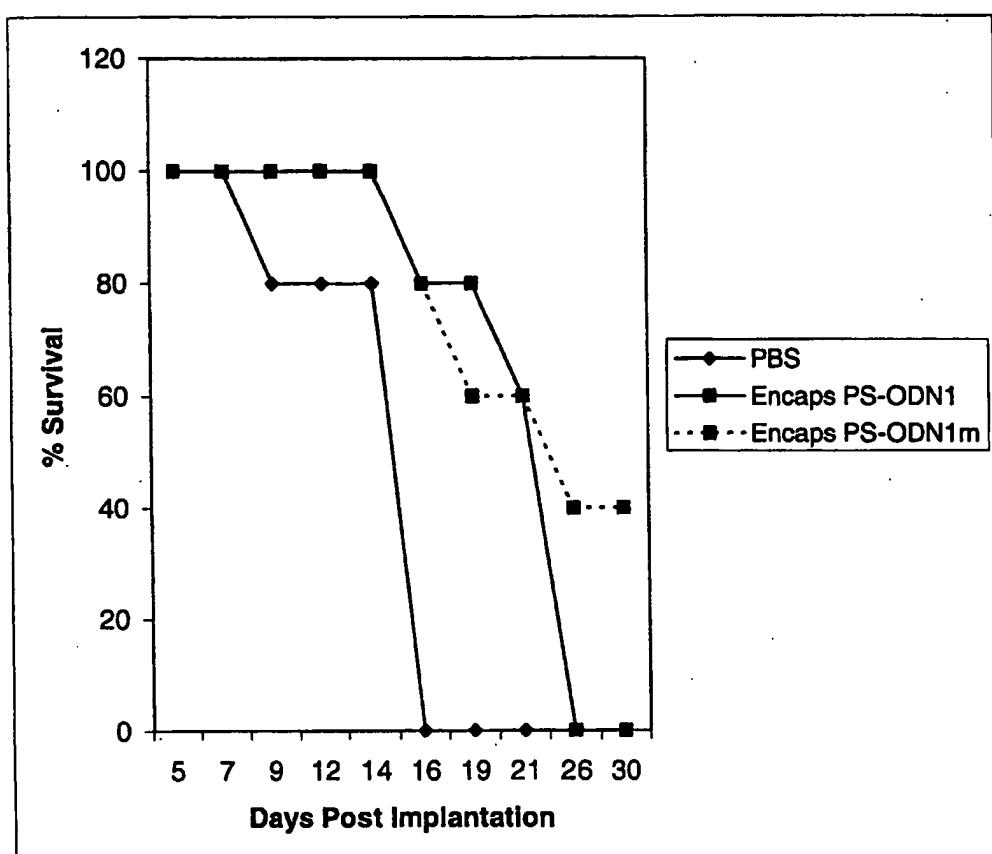


Figure 21

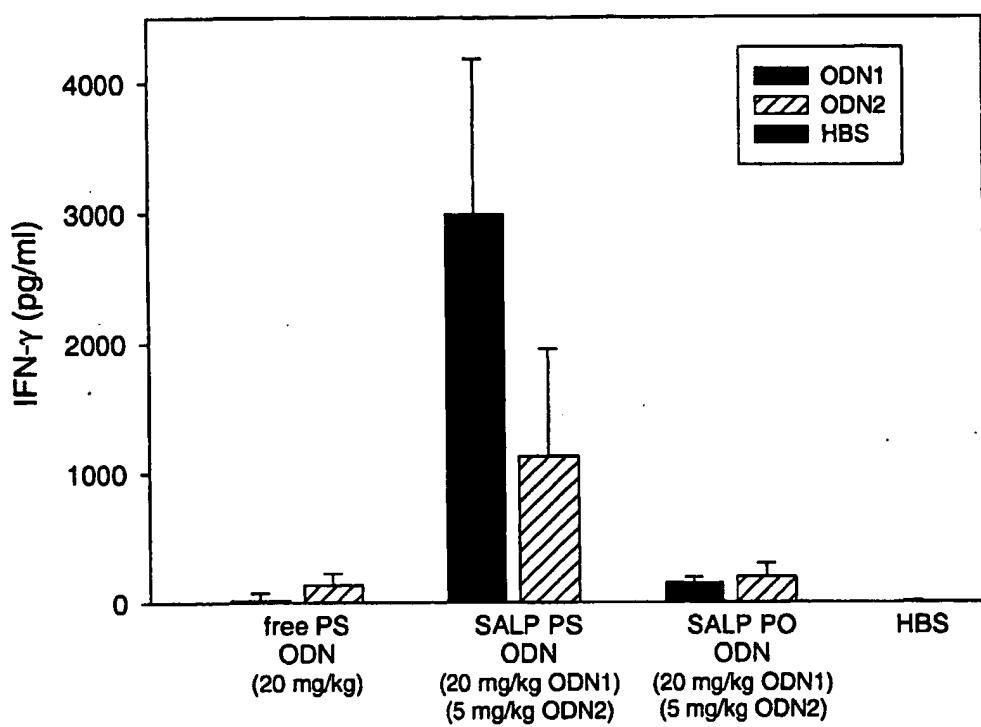


Figure 22

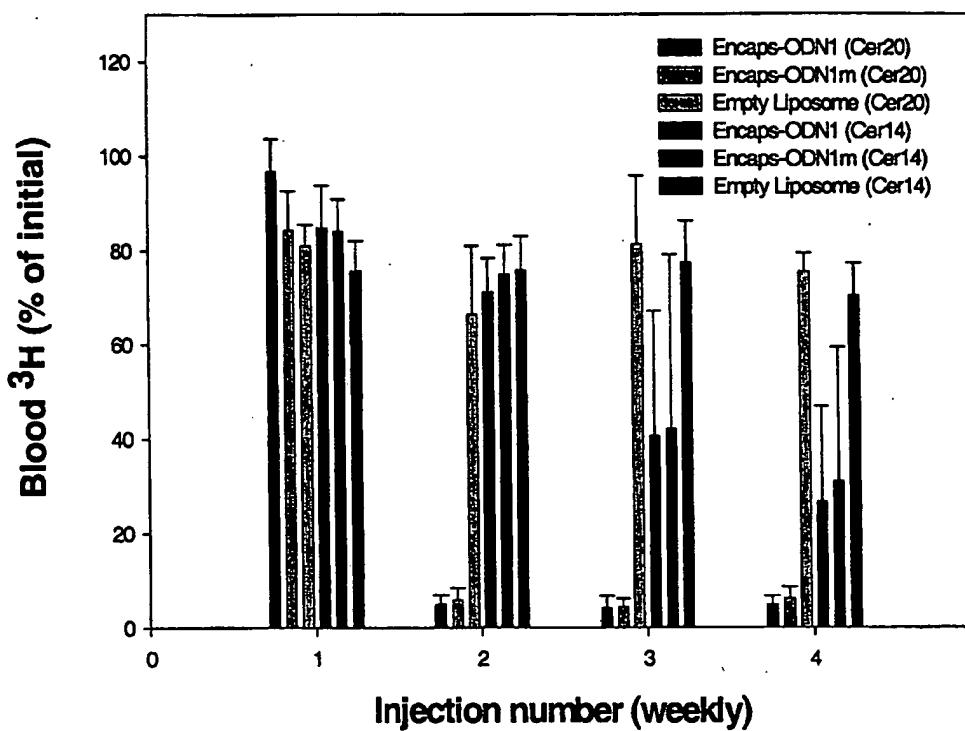


Figure 23

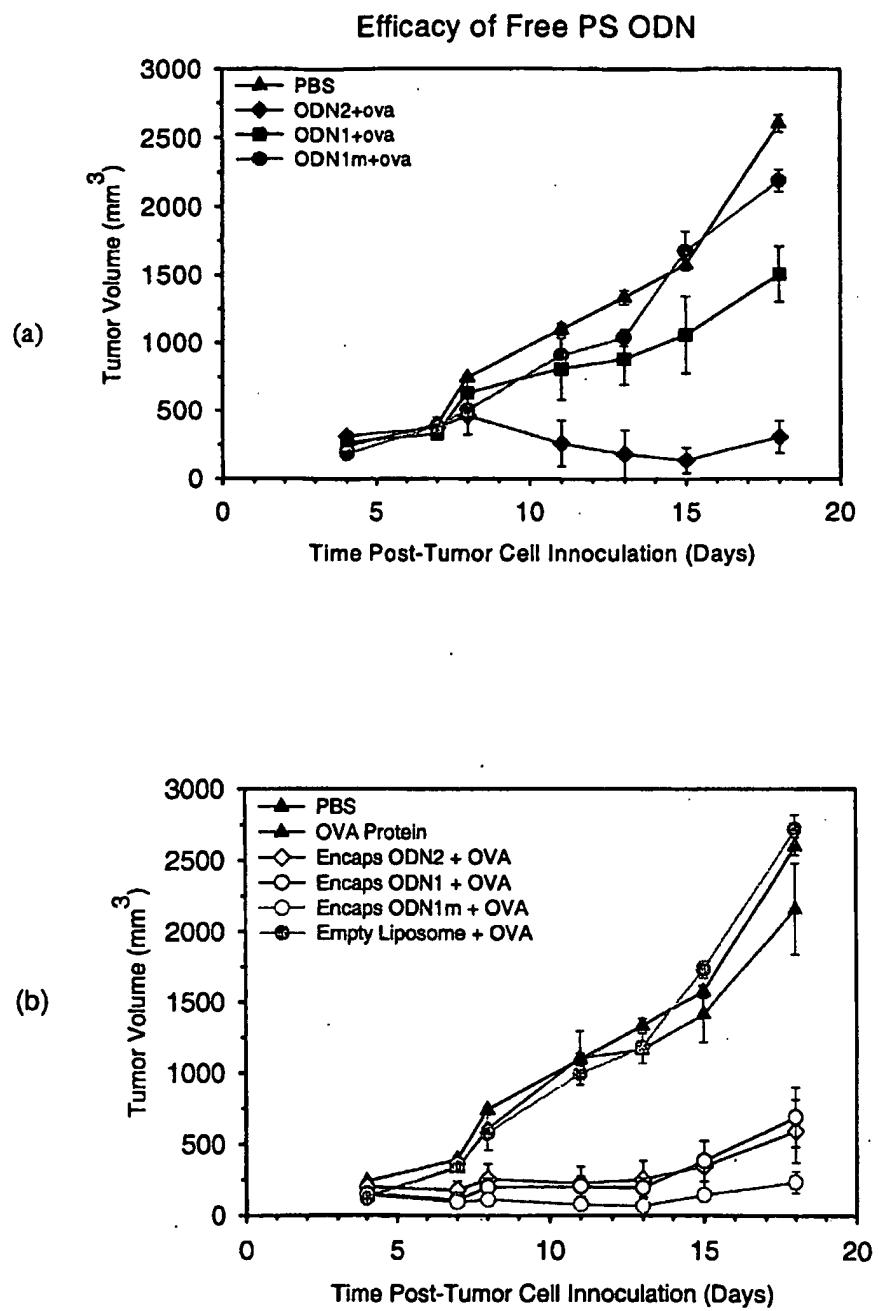


Figure 24

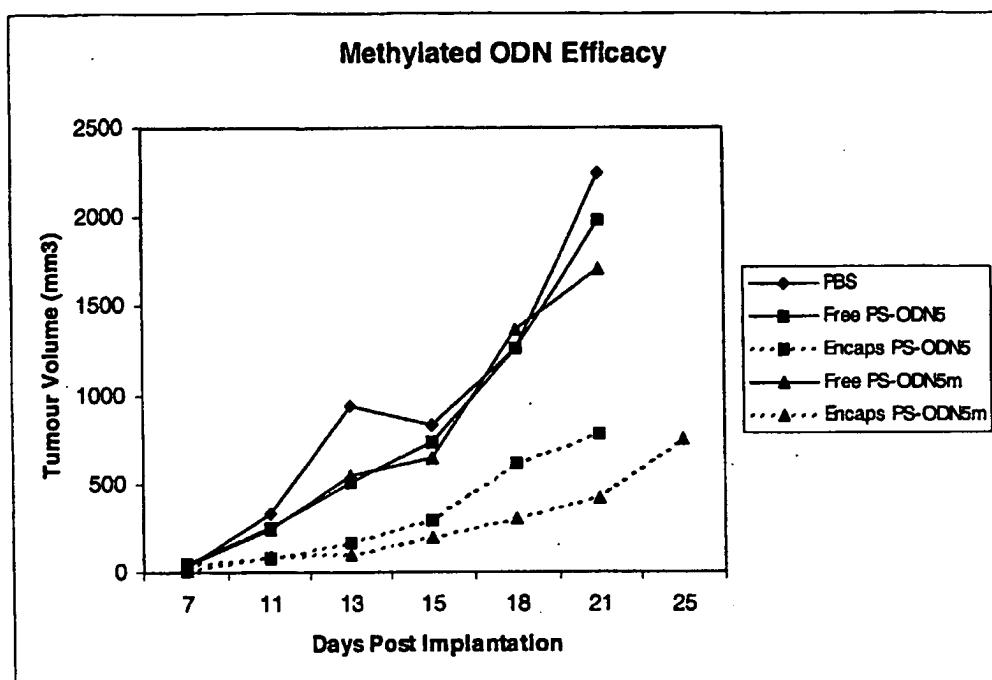


Figure 25

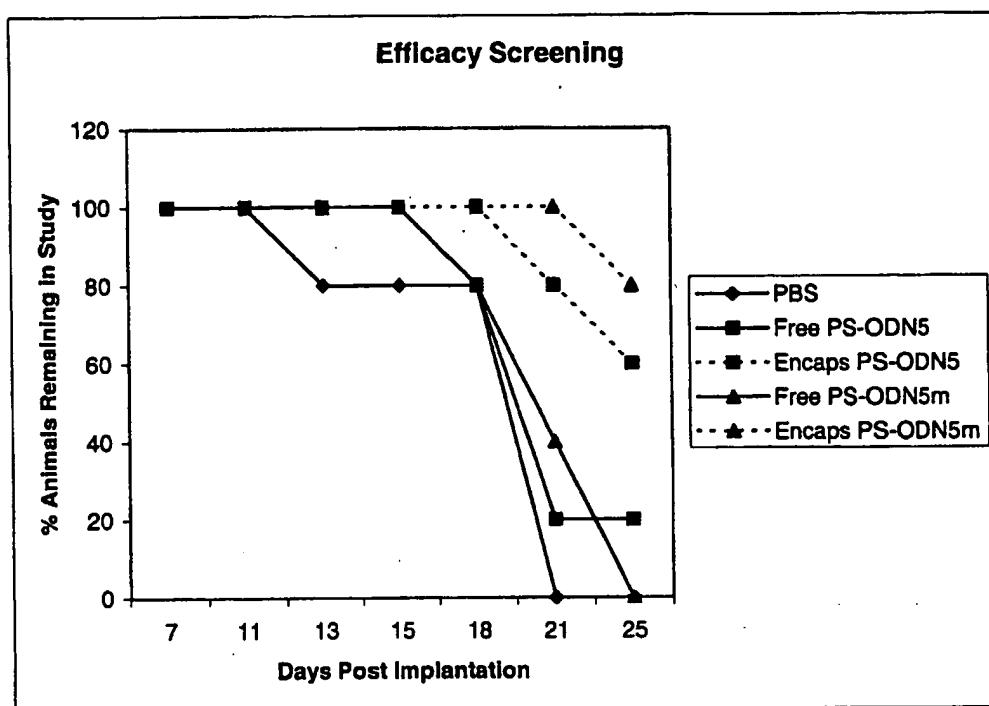


Figure 26

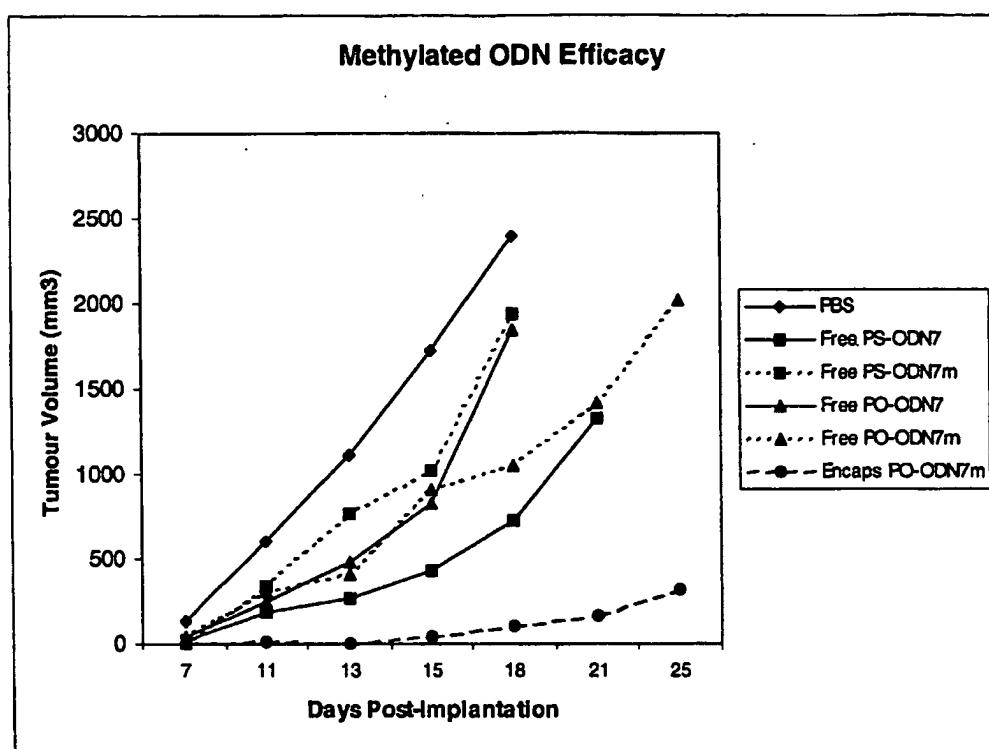


Figure 27

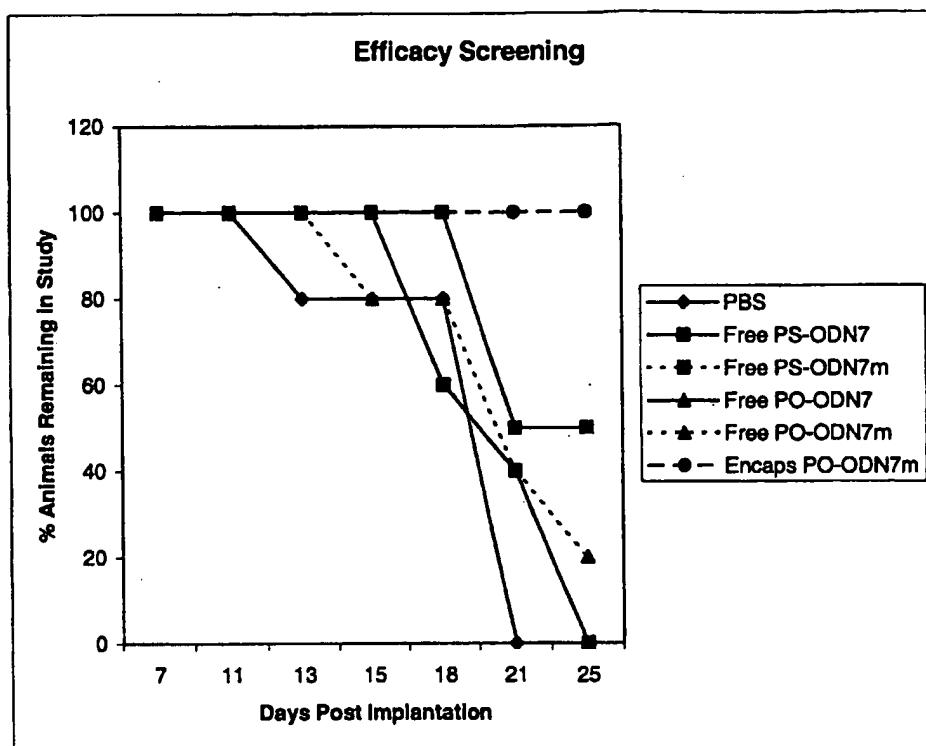


Figure 28

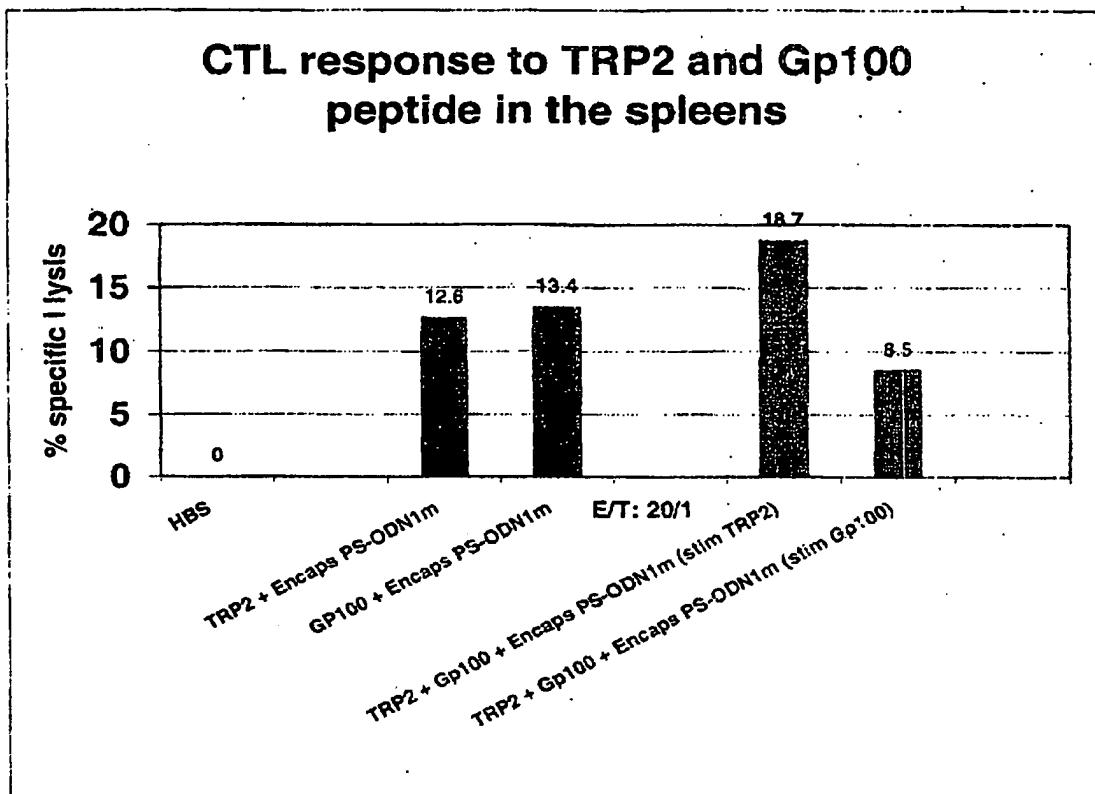


Figure 29

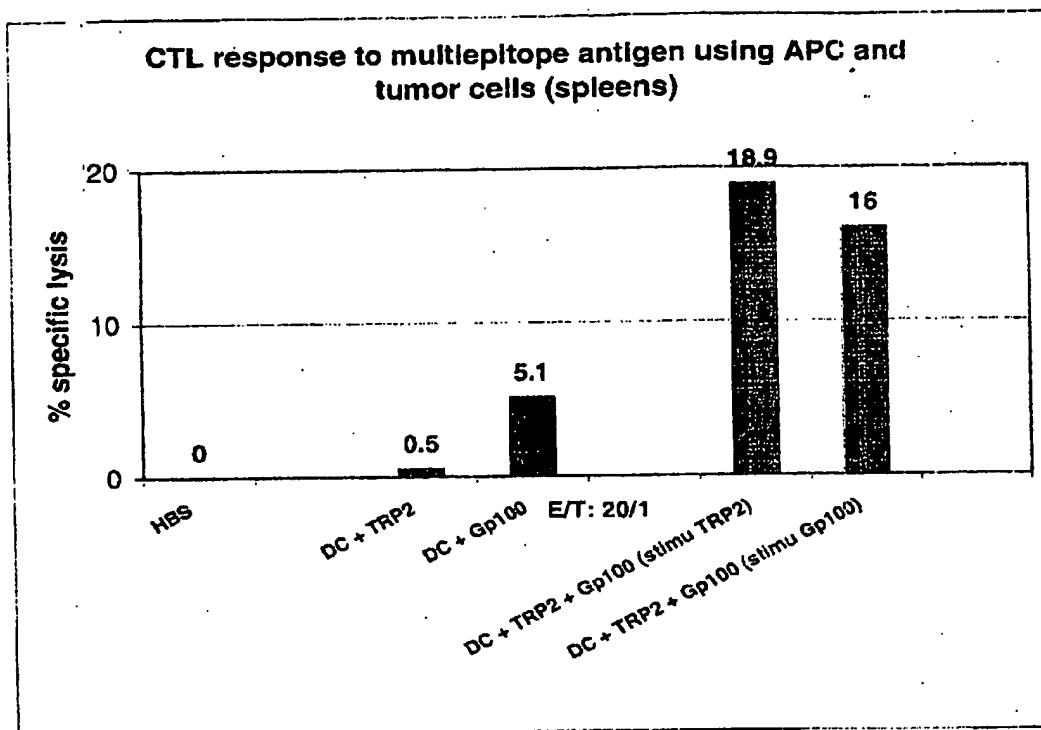


Figure 30

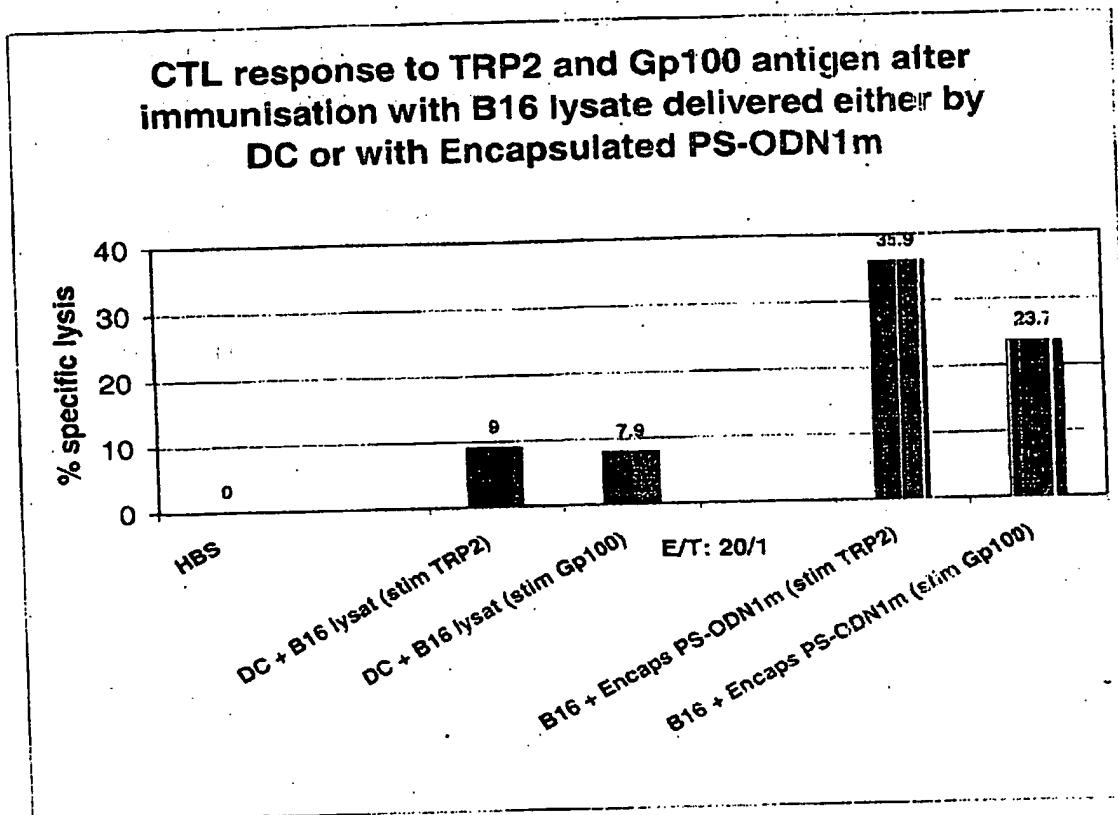


Figure 31